



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 114217

To: Sean McGarry
Location: REM-2C18
Art Unit: 1635
Thursday, February 19, 2004

Case Serial Number: 10/026341

From: Beverly Shears
Location: Remsen Bldg.
RM 1A54
Phone: 571-272-2528

beverly.shears@uspto.gov

Search Notes

SEARCH REQUEST FORM

Requestor's Name: _____ Serial Number: _____
Date: _____ Phone: _____ Art Unit: _____

Search Topic:

Please write a detailed statement of search topic. Describe specifically as possible the subject matter to be searched. Define any terms that may have a special meaning. Give examples or relevant citations, authors, keywords, etc., if known. For sequences, please attach a copy of the sequence. You may include a copy of the broadest and/or most relevant claim(s).

STAFF USE ONLY

Date completed: 02-19-04
Searcher: Bessey C 2528
Terminal time: 20
Elapsed time: _____
CPU time: _____
Total time: 25
Number of Searches: _____
Number of Databases: 1

Search Site

____ STIC
____ CM-1
____ Pre-S

Type of Search

____ N.A. Sequence
____ A.A. Sequence
____ Structure
____ Bibliographic

Vendors

____ IG
____ STN
____ Dialog
____ APS
____ Geninfo
____ SDC
____ DARC/Questel
____ Other CGN

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 11:13:25 ; Search time 1719.07 Seconds
(without alignments)
5592.433 Million cell updates/sec

Title: US-10-026-341A-1

Perfect score: 235

Sequence: 1 agtgcggcagatcatctct.....ccacagttccagacgttga 235

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2888711 seqs, 2045481386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : GenEmbl.*

1: gb_ba.*

2: gb_htg.*

3: gb_in.*

4: gb_om.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pl.*

9: gb_pr.*

10: gb_ro.*

11: gb_sts.*

12: gb_sy.*

13: gb_un.*

14: gb_vi.*

15: em_ba.*

16: em_fun.*

17: em_hum.*

18: em_in.*

19: em_mu.*

20: em_om.*

21: em_or.*

22: em_ov.*

23: em_pat.*

24: em_ph.*

25: em_pl.*

26: em_ro.*

27: em_sts.*

28: em_un.*

29: em_vi.*

30: em_htg_hum.*

31: em_htg_inv.*

32: em_htg_other.*

33: em_htg_mus.*

34: em_htg_pln.*

35: em_htg_rod.*

36: em_htg_mam.*

37: em_htg_vrt.*

38: em_sy.*

39: em_htgo_hum.*

40: em_htgo_mus.*

41: em_htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Match | Length | DB | ID | Description |
|------------|-------|-------|--------|----|-----------|--------------------|
| 1 | 235 | 100.0 | 235 | 6 | AX526280 | Sequence |
| 2 | 235 | 100.0 | 2655 | 9 | HUNTFSF1 | J03133 Human trans |
| 3 | 235 | 100.0 | 2806 | 9 | BC043224 | BC043224 Homo sapi |
| 4 | 235 | 100.0 | 2913 | 9 | AF252284 | AF252284 Homo sapi |
| 5 | 235 | 100.0 | 4300 | 9 | AB039286 | AB039286 Homo sapi |
| 6 | 235 | 100.0 | 73433 | 9 | AC068889 | AC068889 Homo sapi |
| 7 | 222.2 | 94.6 | 2475 | 10 | RNTSP1 | D12768 Rattus norv |
| 8 | 222.2 | 94.6 | 4092 | 10 | AB077988 | AB077988 Rattus no |
| 9 | 222.2 | 94.6 | 206520 | 2 | AC109743 | AC109743 Rattus no |
| 10 | 222.2 | 94.6 | 254686 | 2 | AC097309 | AC097309 Rattus no |
| 11 | 221 | 94.0 | 166697 | 2 | AC021103 | AC021103 Homo sapi |
| 12 | 220.6 | 93.9 | 4128 | 10 | AF022363 | AF022363 Mus muscu |
| 13 | 192.6 | 82.0 | 586 | 11 | G92513 | G92513 S208P6120RE |
| 14 | 192.6 | 82.0 | 3741 | 10 | AF062566 | AF062566 Mus muscu |
| 15 | 192.6 | 82.0 | 138660 | 2 | AC055703 | AC055703 Mus muscu |
| 16 | 192.6 | 82.0 | 205187 | 2 | AC137156 | AC137156 Mus muscu |
| 17 | 84.2 | 35.8 | 2662 | 5 | GGA317960 | AJ317960 Gallus ga |
| 18 | 72.4 | 30.8 | 210390 | 2 | AC110247 | AC110247 Mus muscu |
| 19 | 72.4 | 30.8 | 243137 | 2 | AC119846 | AC119846 Mus muscu |
| 20 | 36.4 | 15.5 | 4954 | 3 | AF213376 | AF213376 Drosophil |
| 21 | 36.4 | 15.5 | 13505 | 2 | AC015161 | AC015161 Drosophil |
| 22 | 36.4 | 15.5 | 54695 | 2 | AC010004 | AC010004 Drosophil |
| 23 | 36.4 | 15.5 | 185497 | 3 | AC010563 | AC010563 Drosophil |
| 24 | 36.4 | 15.5 | 279868 | 3 | AE003518 | AE003518 Drosophil |
| 25 | 36.2 | 15.4 | 225727 | 2 | AC117607 | AC117607 Mus muscu |
| 26 | 35 | 14.9 | 184735 | 9 | AC016180 | AC016180 Homo sapi |
| 27 | 34.6 | 14.7 | 930 | 9 | AY128662 | AY128662 Homo sapi |
| 28 | 34.6 | 14.7 | 1360 | 9 | BC016336 | BC016336 Homo sapi |
| 29 | 34.6 | 14.7 | 1293 | 9 | AB000905 | AB000905 Homo sapi |
| 30 | 34.6 | 14.7 | 89016 | 9 | HS86C11 | AL021807 Human DNA |
| 31 | 34.6 | 14.7 | 141991 | 2 | HS07821D9 | AL121960 Homo sapi |
| 32 | 34.2 | 14.6 | 177806 | 9 | AC093106 | AC093106 Homo sapi |
| 33 | 34.2 | 14.6 | 197246 | 2 | AC118774 | AC118774 Rattus no |
| 34 | 34.2 | 14.6 | 269131 | 2 | AC135469 | AC135469 Mus muscu |
| 35 | 34.2 | 14.6 | 281804 | 2 | AC134869 | AC134869 Mus muscu |
| 36 | 33.8 | 14.4 | 76471 | 9 | AC093743 | AC093743 Homo sapi |
| 37 | 33.8 | 14.4 | 252473 | 2 | AC093972 | AC093972 Rattus no |
| 38 | 33.8 | 14.4 | 271368 | 2 | AC134542 | AC134542 Rattus no |
| 39 | 33.6 | 14.3 | 4745 | 6 | BD174728 | BD174728 Novel iso |
| 40 | 33.4 | 14.2 | 172105 | 2 | AC021197 | AC021197 Homo sapi |
| 41 | 33.4 | 14.2 | 196512 | 9 | AC020911 | AC020911 Homo sapi |
| 42 | 33.2 | 14.1 | 175945 | 2 | AC022147 | AC022147 Homo sapi |
| 43 | 33 | 14.0 | 2544 | 9 | BC006344 | BC006344 Homo sapi |
| 44 | 33 | 14.0 | 2732 | 9 | BC001928 | BC001928 Homo sapi |
| 45 | 33 | 14.0 | 2741 | 9 | BC000425 | BC000425 Homo sapi |

ALIGNMENTS

| | | | | | | |
|------------|--|------------------------------------|--------|-----|--------|-----------------|
| RESULT 1 | AX526280 | Sequence 1 from Patent WO02066071. | 235 bp | DNA | linear | PAT 21-NOV-2002 |
| AX526280 | LOCUS | AX526280 | 235 bp | DNA | linear | PAT 21-NOV-2002 |
| DEFINITION | Sequence 1 from Patent WO02066071. | | | | | |
| ACCESSION | AX526280 | | | | | |
| VERSION | AX526280.1 | GI:25171090 | | | | |
| KEYWORDS | unidentified | | | | | |
| SOURCE | unidentified | | | | | |
| ORGANISM | unclassified. | | | | | |
| REFERENCE | 1 | | | | | |
| AUTHORS | Mauviel A. | | | | | |
| TITLE | Blocking spl transcription factor broadly inhibits extracellular matrix gene expression in vitro and in vivo: implications for the treatments of tissue fibrosis | | | | | |

100

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10973950
2 (bases 1 to 4300)
Takahara,T., Yanagisawa,S. and Akanuma,H.
Direct Submission
Submitted (28-FEB-2000) Hiroshi Akanuma, University of Tokyo,
Graduate School of Arts & Sciences; Komaba 3-8-1, Meguro, Tokyo
153-8902, Japan (E-mail:cakanum@ecc.u-tokyo.ac.jp,
Tel:+81-3-5454-4392(ex.44392))
FEATURES
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BASE COUNT 937 a 1260 c 1016 g 1087 t
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Best Local Similarity 100.0%; Pred. No. 1.1e-63;
Matches 235; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATGGCTGGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGAACAGAGTG 60
DB 2407 ATGGCTGGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGAACAGAGTG 2466
QY 61 GCACAGTACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATCGCAGAGTCTCTGGTG 120
DB 2467 GCACAGTACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATCGCAGAGTCTCTGGTG 2526
QY 121 GGCAGTATGTTGTGGCTCCGCTCCCACTTACAGAACCCAGCAAGTTCTGCAGGACTAC 180
DB 2527 GGCAGTATGTTGTGGCTCCGCTCCCACTTACAGAACCCAGCAAGTTCTGCAGGACTAC 2586
QY 181 CTGGAGTATGCTTAATTCAGTATCATCAAGTAATCCACAGTTCACAGCGTTGA 235
DB 2587 CTGGAGTATGCTTAATTCAGTATCATCAAGTAATCCACAGTTCACAGCGTTGA 2641
RESULT 6
AC068889/c
LOCUS AC068889 73433 bp DNA linear PRI 01-JAN-2003
DEFINITION Homo sapiens 12 BAC RP11-774122 (Roswell Park Cancer Institute
Human BAC Library) complete sequence.
ACCESSION AC068889
VERSION AC068889.35 GI:27452896
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 73433)
Muzny D.M., Adams C., Adio-Ogoola B., Ali-Osman F.R., Allen C.,
Albrooks S.L., Amarantunge H.C., Arc J.R., Ayle M., Banks I.,
Barbaria J., Benton J., Bimage K., Blankenburg K., Bonnin D.,

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Bouck, J., Bowie, S., Brieva, M., Brown, E., Brown, M., Bryant, N.P., Bunay, C., Burch, P., Burkett, C., Burrell, K.L., Byrd, N.C., Carron, T.F., Carter, M., Cavazos, S.R., Chacko, J., Chavez, D., Chen, G., Chen, R., Chiu, D., Chiu, D., Chowdhry, I., Christopoulos, C., Cleveland, C.D., Cox, C., Coyle, M.D., Dathorne, S.R., David, R., Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A., Delaney, K.R., Delgado, O., Denn, A.L., Ding, Y., Dinh, H.H., Douthwaite, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J., Earhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Emerling, S., Escotto, M., Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P., Gabisi, A., Gao, J., Garcia, A., Garner, T., Garza, N., Gill, R., Gorrell, J.H., Guevara, W., Gunaratne, P., Hale, S., Hamilton, K., Han, J., Harris, C., Harris, K., Hart, M., Haviak, P., Hawes, A., Hernandez, J., Hernandez, O., Hodgson, A., Hogues, M., Holloway, C., Hollins, B., Honsi, F., Howard, S., Huber, J., Hulyk, S., Hume, J., Joshikhes, I., Jackson, L.E., Jacobson, B., Jia, Y., Johnson, R., Joliviet, S., Joudah, S., Karlsson, E., Kelly, S., Khan, U., King, L., Korvah, J., Kovar, C., Kratovic, J., Kureshi, A., Landry, N., Leal, B., Lee, Z., Lewis, L.C., Lewis, L., Li, J., Li, Z., Lichtarge, O., Lieu, C., Liu, J., Liu, W., Loulseged, H., Lozano, R.J., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J., Maheshwari, M., Mapua, P., Marandel, I., Martin, R., Martindale, A., Martinez, E., Massey, Z., Mawney, E., McLeod, M.P., Meador, M., Mei, G., Merscher, S., Metzkner, M., Miller, A., Miner, G., Miner, Z., Mitchell, T., Mohabbat, K., Montgomery, K.T., Morgan, M., Morris, S., Moser, M., Neal, D., Nelson, B., Newton, J., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokenwo, S., Ogih, M., Okwuonu, G., Oragunye, N., Ovisdo, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Pickens, R., Primus, E., Pu, L.L., Quiles, M., Ren, Y., Rivers, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Savery, G., Scherer, S., Scott, G., Shen, H., Shim, C., Shooshtari, N., Sisson, I., Sodergren, E., Sonaike, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Svatek, A., Tabot, P., Tamerisa, A., Tamerisa, K., Tang, H., Tansey, J., Taylor, C., Taylor, T., Telford, B., Thomas, R., Thomas, S., Usmami, K., Vasquez, L., Vera, V., Villalon, D., Vinson, R., Wang, Q., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Wallington, S., Williams, G., Williamson, A., Wleczyk, R., Wooden, S., Worley, K., Wu, C., Wu, Y., Wu, Y., Zhou, J., Zorrilla, S., Kucherlapati, R., Weinstein, G. and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 73433)
Worley, K.C.
Direct Submission
Submitted (11-MAY-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 73433)
Worley, K.C.
Direct Submission
Submitted (21-SEP-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 73433)
Worley, K.C.
Direct Submission
Submitted (01-JAN-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Jan 1, 2003 this sequence version replaced gi:23264934.
INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and

SOURCE
ORGANISM
Rattus norvegicus (Norway rat)
Eukaryotes
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sclurognathi; Muridae; Murinae;
Rattus.
1
Takahara, T., Kasahara, D., Mori, D., Yanagisawa, S. and Akanuma, H.
AUTHORS
The trans-spliced variants of Sp1 mRNA in rat
TITLE
Biochem. Biophys. Res. Commun. 298 (1), 155-162 (2002)
MEOLINE
22266831
PUBMED
12379234
2 (bases 1 to 4092)
Mori, D., Takahara, T., Yanagisawa, S. and Akanuma, H.
AUTHORS
Direct Submission
TITLE
Submitted (15-JAN-2002) Hiroshi Akanuma, University of Tokyo,
JOURNAL
Graduate School of Arts & Sciences; Komaba, 3-8-1, Meguro, Tokyo
153-8902, Japan (E-mail: cakanum@ecc.u-tokyo.ac.jp,
Tel: 81-3-5454-3392 (ex. 44392))

```

FEATURES             source
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|-----------------------|-----------------|---|-----------|---------|--------------|
| BASE COUNT | 943 a | 1155 c | 979 g | 1015 t | |
| ORIGIN | | | | | |
| Query Match | 94.6%; | Score | 222.2; | DB 10; | Length 4092; |
| Best Local Similarity | 96.6%; | Pred. No. | 1.3e-59; | | |
| Matches 227; | Conservative 0; | Mismatches 8; | Indels 0; | Gaps 0; | |
| QY | 1 | ATGGCTGGCAGATCATCTCTTCTCTCTCTGGGGCTACCCCTACCTCAAAGGAACAGATG | 60 | | |
| Db | 2263 | ATGGCTGGCAGATCATCTCTTCTCTCTCTGGGGCTACCCCTACCTCAAAGGAACAGATG | 2322 | | |
| QY | 61 | GCAGCAGTACCAATGGCGAGCAATGGCAGTCAGTCTTCCAAAGAAATGCACAGTCTCTGTG | 120 | | |
| Db | 2323 | GCACAGTACCAATGGCGAGCAATGGCAGTCAGTCTTCCAAAGAACGCACAGTCTCTGTG | 2382 | | |
| QY | 121 | GCAGTATGTTGTGGCTGCCGCTCCCACTTACAGAACCGCAGTTCGACAGACTAC | 180 | | |
| Db | 2383 | GGCAGTATGTTGTGGCTGCTACCCCCAACTTACAGAACCGCAGTTCGACAGCTCTAC | 2442 | | |
| QY | 181 | CTGGAGTGTAGTCCTAAATATTCAGTATCCAGTATCCACAGTTCACAGCCGTGA | 235 | | |
| Db | 2443 | CTGGAGTATGCTTAATATTCAGTATCCAGTATCCACAGTTCACAGCTTGA | 2497 | | |

| | | | | | |
|------------|---|-----------|-----|--------|-----------------|
| RESULT 9 | AC109743 | 206520 bp | DNA | linear | HTG 19-NOV-2005 |
| LOCUS | AC109743 | | | | |
| DEFINITION | Rattus norvegicus clone CH230-65D10, *** SEQUENCING IN PROGRESS | | | | |
| ACCESSION | ***, 3 unordered pieces. | | | | |
| | AC109743 | | | | |

VERSION

AC109743.5 GI:25072885

KEYWORDS

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.

SOURCE

Rattus norvegicus (Norway rat)

ORGANISM

Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE

1 (bases 1 to 206520)

AUTHORS

Munzy D. Marie, Metsker M. Lee, Abramson S., Adams C., Alder J.,
Allen C., Allen H., Albrooks S., Amin A., Anguiano D.,
Anyalebechi V., Aoyagi A., Ayodeji M., Baca E., Baden H.,
Baldwin D., Bandaranaike D., Barber M., Barnstead M., Benahmed F.,
Biswal K., Blair J., Blankenburg K., Blyth P., Brown M.,
Bryant N., Buhay C., Burch P., Burrell K., Calderon E.,
Cardenas V., Carter K., Cavazos J., Ceasar H., Chen A.,
Chacko J., Chavez D., Chen R., Chen Y., Chen Z., Chu J.,
Claveland C., Cockrell R., Cox C., Coyle M., Cree A., D'Souza L.,
Davila M., Davis C., Davy-Carroll L., De Anda C., Dederich D.,
Delgado O., Denson S., Deramo C., Ding Y., Dinh H., Divya K.,
Draper H., Dugan-Rocha S., Dunn A., Durbin K., Duval B., Eaves K.,
Egan A., Escotto M., Eugene C., Evans C.A., Falls T., Fan G.,
Fernandez S., Finley M., Flagg N., Forbes L., Foster M., Foster P.,
Fraser C.M., Gabisi A., Ganta R., Garcia A., Garner F., Garza M.,
Gregoire G., Geer K., Gill R., Grady M., Guerra W., Guevara W.,
Gunaratne P., Haaland W., Hamill C., Hamilton C., Hamilton K.,
Harvey Y., Havlak P., Hawes A., Henderson N., Hernandez J.,
Hernandez R., Hines S., Hladun S.L., Hodgson A., Hogues M.,
Hollins B., Howells S., Hulyk S., Hume J., Idlebird D., Jackson A.,
Jackson L., Jacob L., Jiang H., Johnson B., Johnson R., Jolivet A.,
Karpach S., Kelly S., Khan Z., King L., Kovar C.,
Kowis C., Kraft C.L., Lebow H., Levan J., Lewis L., Li Z., Liu J.,
Liu J., Liu W., Liu Y., London P., Longacre S., Lopez J.,
Lorensuwa L., Loulseghe H., Lozado R.J., Lu X., Ma J.,
Maheshwari M., Mahindratne M., Mahmoud M., Malloy K., Mangum A.,
Mangum B., Mapua P., Martin K., Martin R., Martinez E.,
Mawney S., McLeod M.P., McNeill T.Z., Meenen E.,
Milosavljevic A., Miner G., Minja E., Montemayor J., Moore S.,
Morgan M., Morris K., Morris S., Munidasa M., Murphy M., Nair L.,
Nankervis C., Neal D., Newton N., Nguyen N., Norris S.,
Nwaekemah O., Okwuon G., Olarnpunsagoon A., Pal S., Parks K.,
Pasternak S., Paul H., Perez A., Perez L., Pfannkuch C.,
Plopper F., Poindexter A., Popovic D., Primus E., Pu L.,
Puzo M., Quiroz J., Rachlin E., Reeves K., Regier M.A., Reigh R.,
Reilly B., Reilly M., Ren Y., Reuter M., Richards S., Riggs F.,
Rives C., Rodkey T., Rojas A., Rose M., Rose R., Ruiz S.J.,
Sanders M., Savery G., Scherer S., Scott G., Shatsman S., Shen H.,
Shetty J., Shvartsbeyn A., Sison I., Sitter C.D., Smajls D.,
Sneed A., Sodergren E., Song X.-Z., Sorrelle R., Sosa J.,
Steimle M., Strong R., Sutton A., Svatek A., Tabor P., Taylor C.,
Taylor T., Thomas N., Thomas S., Tingey A., Trejos P., Usmani K.,
Valas R., Vera V., Villanasa D., Waldron L., Walker B., Wang J.,
Wang Q., Wang S., Warren J., Warren R., Wei X., White F.,
Williams G., Willson R., Mleczyk R., Wooden H., Worley K.,
Wright D., Wright R., Wu J., Yakub S., Yen J., Yoon L., Yoon V.,
Yu F., Zhang J., Zhou X., Zhou X., Zhao S., Dunn D., von
Niederhausern A., Weiss R., Smith D.R., Holt R.A., Smith H.O.,
Weinstock G. and Gibbs R.A.

TITLE

Unpublished

JOURNAL

2 (bases 1 to 206520)

AUTHORS

Worley K.C.

TITLE

Direct Submission

JOURNAL

Submitted (07-FEB-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

REFERENCE

3 (bases 1 to 206520)

AUTHORS

Rat Genome Sequencing Consortium.

TITLE

Direct Submission

JOURNAL

Submitted (19-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

COMMENT

On Nov 19, 2002 this sequence version replaced gi:23195412.
The sequence in this assembly is a combination of BAC based reads

and whole genome shotgun sequencing reads assembled using Atlas
(<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: GQNU

Center clone name: CH230-65D10

----- Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 168784 bases at least Q40

Consensus quality: 173457 bases at least Q30

Consensus quality: 175960 bases at least Q20

Estimated insert size: 178632; sum-of-contigs estimation

Quality coverage: 5x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length

* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 3 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1 76730 contig of 76730 bp in length

* 76731 76830: gap of unknown length

* 76831 204784: contig of 127954 bp in length

* 204785 204884: gap of unknown length

* 204885 206520: contig of 1636 bp in length.

----- Location/Qualifiers

1. 206520

/organism="Rattus norvegicus"

/mol_type="genomic DNA"

/db_xref="taxon:10116"

/clone="CH230-65D10"

101270..102941

/note="wgs contig"

168407..170678

/note="wgs contig"

BASE COUNT 49558 a 37786 c 38536 g 52117 t 28523 others

ORIGIN

Query Match 94.6%; Score 222.2; DB 2; Length 206520;

Best Local Similarity 96.6%; Pred. No. 1.9e-59;

Matches 227; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 1 ATGGCTGGCAGATCATCTCTCTCTCTCTGGGGCTACCCCTACCTCAAGGGAACAGAGTG 60

Db 14467 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGGAACAGAGTG 14526

QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATCGCAGTCTCTGGTG 120

Db 14527 GCACAGTACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATCGCAGTCTCTGGTG 14586

QY 121 GGCAGTATGTGTGGCTGGCTGCCGTCCCAACTTACAGAACCCAGCAAGTCTGACGAGCTAC 180

Db 14587 GGCAGTATGTGTGGCTGGCTGCCGTCCCAACTTACAGAACCCAGCAAGTCTGACGAGCTAC 14646

QY 181 CTGAGTGTATGCTTAATATTCAGTATCAAGTAAATCCACAGTTCAGACCGCTTGA 235

Db 14647 CTGGAGTAATGCTAATATTCAGTATCAATCAAGTAATCCACAGTTCACAGACTGTGA 14701

RESULT 10
AC097309
LOCUS
DEFINITION Rattus norvegicus clone CH230-97M10, WORKING DRAFT SEQUENCE, 2
unordered pieces.

AC097309 254686 bp DNA linear HTG 10-MAY-2003
AC097309.6 GI:30521315
HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
Rattus norvegicus (Norway rat)
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE 1 (bases 1 to 254686)
Muzny,D,Marie., Metzker,M.Lee., Abramson,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Biswal,K., Blair,J., Blakenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
Chacko,J., Chavez,D., Chen,G., Chen,K., Chen,Y., Chen,Z., Chu,J.,
Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Evans,C., Evans,C.A., Falls,T., Fan,G.,
Fernandes,S., Finley,M., Flegg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
Gebregeorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
Gunaratne,P., Haaland,W., Hamil,C., Hamilton,C., Hamilton,K.,
Harvey,Y., Havlak,P., Hawes,A., Henderson,N., Hernandez,J.,
Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hoques,M.,
Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A.,
Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpachy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,F., Longacre,S., Lopez,J.,
Lorenshaw,L., Loulseghe,H., Lozado,R.J., Lu,X., Ma,J.,
Maheshwari,M., Mahindartne,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
Mawhinney,S., McLeod,M.P., McNeill,I.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nait,L.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,
Nwakaleleh,O., Okwuonu,G., Olarnpunsagoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfannkuch,C.,
Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L.,
Puazo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,W.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,W., Savery,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,J., Shvartabeyn,A., Sisson,I., Sitter,C.D., Smajd,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K.,
Vallas,R., Vera,V., Villalana,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wleczyk,R., Wooden,H., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,X., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.

Direct Submission
Unpublished
2 (bases 1 to 254686)
Worley,K.C.
Direct Submission
Submitted (14-OCT-2001) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 254686)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (10-MAY-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On May 10, 2003 this sequence version replaced gi:23268851.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

Center: Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
Project Information
Center project name: GIGO
Center clone name: CH230-97M10
Summary Statistics
Assembly program: Atlas 3.0;
Consensus quality: 240288 bases at least Q40
Consensus quality: 242808 bases at least Q30
Consensus quality: 244806 bases at least Q20
Estimated insert size: 254704; sum-of-contigs estimation
Quality coverage: 9x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)
* NOTE: This sequence may represent more than one 'clone'
* NOTE: This is a 'working draft' sequence. It currently
consists of 2 contigs. The true order of the pieces
is not known and their order in this sequence record is
arbitrary. Gaps between the contigs are represented as
runs of N, but the exact sizes of the gaps are unknown.
This record will be updated with the finished sequence
as soon as it is available and the accession number will
be preserved.

* 1 253477: contig of 253477 bp in length
* 253478 253577: gap of unknown length
* 253578 254686: contig of 1109 bp in length.

Location/Qualifiers
1..254686
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-97M10"
1..3186
/note="wgs contig"
251549..252396
/note="clone boundary
clone_end:Sp6
site:EcoRI
end sequence:BH283956"
BASE COUNT 63545 a 59875 c 59227 g 63989 t 9050 others
ORIGIN

Query Match 94.6%; Score 222.2; DB 2; Length 254686;
Best Local Similarity 96.6%; Pred. No. 1.9e-59;
Matches 227; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 1 ATGGCTGGCAGATCATCTTCCTCTCCCTGCGGTACCCCTACCTCAAGGACAGAGTG 60
|||||
Db 212410 ATGGCTGGCAGATCATCTTCCTCTCCCTGCGGTACCCCTACCTCAAGGACAGAGTG 212469
|||||


```
OY 61 GCAGAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAAAGATCCACAGTCTCTGGTG 120
Db 212470 GCAACAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAAAGATCCACAGTCTCTGGTG 212529
OY 121 GCAGATGATCTGGTGGTGGTCCCACTTACAGAACCCAGCAAGTCTTGACAGCACTAC 180
Db 212530 GCAGATGATCTGGTGGTGGTACCCCACTTACAGAACCCAGCAAGTCTTGACAGCACTAC 212589
OY 181 CTGGAGTGTGCTTAATATTCAGTATCAAGTAAATCCACAGTTCACAGCCGTGA 235
Db 212590 CTGGAGTGTGCTTAATATTCAGTATCAAGTAAATCCACAGTTCACAGCCGTGA 212644

RESULT 11
AC021103 166697 bp DNA linear HTG 07-JUL-2000
LOCUS Homo sapiens chromosome UL clone RP11-147A18, WORKING DRAFT
DEFINITION SEQUENCE. 43 unordered pieces.
ACCESSION AC021103
VERSION AC021103.7 GI:8099088
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 166697)
AUTHORS Waterston,R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 166697)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (14-JAN-2000) Genome Sequencing Center, Washington
University School of Medicine, 444 Forest Park Parkway, St. Louis,
MO 63108, USA
COMMENT On May 26, 2000 this sequence version replaced gi:8018258.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H_NEO147A18
----- Summary Statistics -----
Sequencing vector: M13; 66%
Sequencing vector: Plasmid; 34%
Chemistry: Dye-primer ET; 66% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 151987 bases at least Q40
Consensus quality: 156240 bases at least Q30
Consensus quality: 158337 bases at least Q20
Insert size: 184000; agarose-fp
Insert size: 162497; sum-of-contigs
Quality coverage: 3.88 in Q20 bases; agarose-fp
Quality coverage: 4.39 in Q20 bases; sum-of-contigs
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 43 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 1466: contig of 1466 bp in length
* 1467 1566: gap of unknown length
* 1567 2896: contig of 1330 bp in length
* 2897 2996: gap of unknown length
* 2997 4016: contig of 1020 bp in length
* 4017 4116: gap of unknown length
* 4117 5406: contig of 1290 bp in length

5506: gap of unknown length
5507 6642: contig of 1136 bp in length
6643 8257: gap of unknown length
8258 8357: gap of unknown length
8358 10742: contig of 2385 bp in length
10743 10842: gap of unknown length
10843 12190: contig of 1348 bp in length
12191 12290: gap of unknown length
12291 13569: contig of 1279 bp in length
13570 15855: gap of unknown length
15856 17977: contig of 2186 bp in length
17978 18078: gap of unknown length
18079 20177: contig of 2022 bp in length
20178 22120: gap of unknown length
22121 24582: contig of 2100 bp in length
24583 24682: gap of unknown length
24683 27320: contig of 2638 bp in length
27321 29908: gap of unknown length
29909 30088: contig of 2488 bp in length
30089 32534: gap of unknown length
32535 35494: contig of 2526 bp in length
35495 37552: gap of unknown length
37553 41357: contig of 2859 bp in length
41358 43778: gap of unknown length
43779 47604: contig of 2321 bp in length
47605 50117: gap of unknown length
50118 53652: contig of 3626 bp in length
53653 56657: gap of unknown length
56658 60073: contig of 2905 bp in length
60074 60174: gap of unknown length
60175 64263: contig of 3316 bp in length
64264 68859: contig of 4090 bp in length
68860 72754: gap of unknown length
72755 77167: contig of 4496 bp in length
77168 81554: gap of unknown length
81555 84717: contig of 4287 bp in length
84718 89925: gap of unknown length
89926 94331: contig of 3062 bp in length
94332 98565: gap of unknown length
98566 105959: contig of 4133 bp in length
105960 112344: gap of unknown length
112345 119806: contig of 7295 bp in length
119807 127259: gap of unknown length
127260 136297: contig of 6285 bp in length
136298 136397: gap of unknown length
```



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source 1. 586
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="129S1/SvImJ"
/db_xref="taxon:10090"
/map="+" 15 22-551 103388565-103389094"
/clone_lib="129S1/SvImJ"
<1. >586
STS
BASE COUNT 141 a 159 c 138 g 148 t
ORIGIN
Query Match 82.0%; Score 192.6; DB 11; Length 586;
Best Local Similarity 92.3%; Pred. No. 3e-50;
Matches 217; Conservative 0; Mismatches 9; Indels 9; Gaps 1;

QY 1 ATGGCTGGCAGATCATCTCTCTCTCTCTGGGGCTACCCCTCAAGGACAGAGTG 60
Db 333 ATGGCTGGCAGATCATCTCTCTCTCTCTGGGGCTACCCCTCAAGGACAGAGTG 392

QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGAGTCTTCCAGAAATCGCAGTCTCTGGTG 120
Db 393 GCACAGTACCAATGGCAGC-----GAGTCTTCCAGAACCCGACAGTCTCTGGTG 443

QY 121 GCAGATGTTGTGGCTGCGGCTCCCACTTACAGAACCCAGCAAGTCTTCAAGCACTAC 180
Db 444 GCAGATGTTGTGGCTGCTACCCCACTTACAGAACCCAGCAAGTCTTCAAGCACTCC 503

QY 181 CTGAGTGATGCTTAATATTCAGTATCAAGTAAATCCACAGTTCACAGCCGTTGA 235
Db 504 CTGAGTATGCTTAATATTCAGTATCAAGTAAATCCACAGTTCACAGCTGTGA 558

RESULT 14
AF062566 3741 bp mRNA linear ROD 22-JUN-2001
LOCUS Mus musculus transcription factor Spl mRNA, complete cds.
ACCESSION AF062566
VERSION AF062566.1 GI:3135322
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 3741)
Yajima, S., Lee, S.H., Minowa, T. and Mouradian, M.M.
Sp family transcription factors regulate expression of rat D2
dopamine receptor gene
DNA Cell Biol. 17 (5), 471-479 (1998)
JOURNAL 98290554
MEDLINE 9828590
PUBMED
REFERENCE 2 (bases 1 to 3741)
Yajima, S., Lee, S.-H., Minowa, T. and Mouradian, M.M.
Direct Submission
Submitted (24-APR-1998) GPU/ETB, NINDS/NIH, 10 Center Dr MSC1406,
Bethesda, MD 20892-1406, USA
JOURNAL Location/Qualifiers
FEATURES
source
1. 3741
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/cell_line="NB41A3"
/tissue_type="neuroblastoma"
37..2382
/codon_start=1
/product="transcription factor Spl"
/protein_id="AAC16484.1"
/db_xref="GI:3135323"
translation="MSDDHSDVEYAVVKIEKVDGNGNGSGGGAFAFSQTSST
GSSSSGGGGSGSPSLAALAACTSRLESNNNSQSPSGGGLDLTAQVL
SQGANQWQIISSSGATPFSKSGNSTNGSSNNRVSGQIVYVATPFLNQCVL
TGLPWNPIQVQIPQFQVDDQLQFAAQVQDQGGQIQIPGANQIIPNRG
SGNNIIPNLQQVAPLQGLANNVLSCQTVVNPVAVLNGNITLLPVNSAATL
TPSSQAGTISSSGQESSQPVTSCTAISASLSVSSQSSSFFFTNANSYSTTTTNS
-----Genome Center
Center: Harvard Partners Genome Center
Center Code: HPGC
Web site: http://www.hpcgg.org/sequence/mouse.html
Contact: hpcg@medel.mgh.harvard.edu
-----Summary Statistics
```

```
Center project name: AAS
Sequencing vector: pUC18; 108752
Chemistry: Dye-terminator Big Dye; 100%
*Consensus quality: 126666 at least Q20
*Consensus quality: 122952 at least Q30
*Consensus quality: 116499 at least Q40
Estimated insert size: agarose-PP - N/A
**Estimated insert size: 138140 - sum-of-contigs
Quality coverage: agarose-PP - N/A
Quality coverage: 9.1 x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 37 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
1 10740: contig of 10740 bp in length
* 10741 10760: gap of unknown length
* 10761 18887: contig of 8127 bp in length
* 18888 18907: gap of unknown length
* 18908 26674: contig of 7767 bp in length
* 26675 26694: gap of unknown length
* 26695 31485: contig of 4791 bp in length
* 31486 31505: gap of unknown length
* 31506 36220: contig of 4715 bp in length
* 36221 36240: gap of unknown length
* 36241 43190: contig of 6950 bp in length
* 43191 43210: gap of unknown length
* 43211 47455: contig of 4245 bp in length
* 47456 47475: gap of unknown length
* 47476 52355: contig of 4880 bp in length
* 52356 52375: gap of unknown length
* 52376 57154: contig of 4779 bp in length
* 57155 57174: gap of unknown length
* 57175 63909: contig of 6735 bp in length
* 63910 63929: gap of unknown length
* 63930 70719: contig of 6790 bp in length
* 70720 70739: gap of unknown length
* 70740 73652: contig of 2913 bp in length
* 73653 73672: gap of unknown length
* 73673 79581: contig of 5909 bp in length
* 79582 83557: contig of 3956 bp in length
* 83558 83577: gap of unknown length
* 83578 89230: contig of 5853 bp in length
* 89231 89250: gap of unknown length
* 89251 93443: contig of 4193 bp in length
* 93444 93464: gap of unknown length
* 93464 97169: contig of 3706 bp in length
* 97170 97189: gap of unknown length
* 97190 102251: contig of 5062 bp in length
* 102252 102271: gap of unknown length
* 102272 107091: contig of 4820 bp in length
* 107092 107111: gap of unknown length
* 107112 110480: contig of 3369 bp in length
* 110481 110500: gap of unknown length
* 110501 113418: contig of 2918 bp in length
* 113419 113438: gap of unknown length
* 113439 116688: contig of 3230 bp in length
* 116689 118403: contig of 1715 bp in length
* 118404 118423: gap of unknown length
* 118424 120544: contig of 2121 bp in length
* 120545 120564: gap of unknown length
* 120565 121933: contig of 1429 bp in length
* 121994 122013: gap of unknown length
* 122014 122930: contig of 917 bp in length
* 122931 122950: gap of unknown length
* 122951 124509: contig of 1559 bp in length
* 124510 124529: gap of unknown length
*
124530 126851: contig of 2322 bp in length
* 126852 128871: gap of unknown length
* 128872 128496: contig of 1625 bp in length
* 128497 128516: gap of unknown length
* 128517 129949: contig of 1433 bp in length
* 129950 129969: gap of unknown length
* 129970 132003: contig of 2034 bp in length
* 132004 132023: gap of unknown length
* 132024 133605: contig of 1582 bp in length
* 133606 133625: gap of unknown length
* 133626 134100: contig of 475 bp in length
* 134101 134120: gap of unknown length
* 134121 135517: contig of 1397 bp in length
* 135518 135537: gap of unknown length
* 135538 136468: contig of 931 bp in length
* 136469 136488: gap of unknown length
* 136489 137236: contig of 748 bp in length
* 137237 137256: gap of unknown length
* 137257 138860: contig of 1604 bp in length.
*
FEATURES
Source
1. 138860
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL6/J"
/db_xref="taxon:10090"
/clone="RP23-399N14"
/sex="male"
1. 10740
/note="assembly_name:Contig270"
10761..18887
/note="assembly_name:Contig269"
18908..26674
/note="assembly_name:Contig268"
26695..31485
/note="assembly_name:Contig267"
31506..36220
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36241..43190
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43211..47455
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52376..57154
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70740..73652
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73673..79581
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79602..83557
/note="assembly_name:Contig257"
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89251..93443
/note="assembly_name:Contig255"
93464..97169
/note="assembly_name:Contig254"
97190..102251
/note="assembly_name:Contig253"
102272..107091
/note="assembly_name:Contig252"
107112..110480
/note="assembly_name:Contig251"
110501..113418
/note="assembly_name:Contig250"
113439..116688
/note="assembly_name:Contig249"
116689..118403
/note="assembly_name:Contig248"
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misc_feature 118424..120544
/notes="assembly_name:Contig247"
misc_feature 120565..121993
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clone_end:SP6
vector_side:left"
misc_feature 122014..122930
/notes="assembly_name:Contig245"
misc_feature 122951..124509
/notes="assembly_name:Contig244"
misc_feature 124530..126851
/notes="assembly_name:Contig243"
misc_feature 126872..128496
/notes="assembly_name:Contig242"
misc_feature 128517..129949

Query Match      82.0%; Score 192.6; DB 2; Length 138860;
Best Local Similarity 92.3%; Pred. No. 5e-50;
Matches 217; Conservative 0; Mismatches 9; Indels 9; Gaps 1;

QY 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
Db 108054 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 107995

QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGAAGTCTTCCAAAGATCGCACAGTCTCTGGTG 120
Db 107994 GCAACAGTACCAATGGCAGC-----GAGTCTTCCAAAGAACCGCACAGTCTCTGGTG 107944

QY 121 GGCAGTATGTTGGTGCGCTCCCACTTACAGAACGAGAGTTCTGACAGGACTAC 180
Db 107943 GGCAGTATGTTGGTGCTGTACCCCACTTACAGAACGAGAGTTCTGACAGGCTCC 107884

QY 181 CTGGAGTATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGCCGTTGA 235
Db 107883 CTGGAGTATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGCTGTTGA 107829

```

Search completed: February 18, 2004, 15:26:04
Job time : 1724.07 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 08:17:00 ; Search time 241.401 Seconds
(without alignments)
2627.862 Million cell updates/sec

Title: US-10-026-341A-1
Perfect score: 235
Sequence: 1 agcgctgcagatcatctct.....ccacagttccagaccgttga 235

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2552756 seqs, 1349719017 residues
Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

- Database : N Geneseq 19Jun03.*
- 1: /SIDS1/gcgdata/geneseq/geneseq-emb1/NA1980.DAT.*
 - 2: /SIDS1/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
 - 3: /SIDS1/gcgdata/geneseq/geneseq-emb1/NA1982.DAT.*
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 - 22: /SIDS1/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
 - 23: /SIDS1/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----|--------------------|
| 1 | 235 | 100.0 | 235 | 24 | Spl gene fragment. |
| 2 | 235 | 100.0 | 2862 | 25 | Human nucleic acid |
| 3 | 235 | 100.0 | 3090 | 22 | Human polynucleoti |
| 4 | 36.4 | 15.5 | 2691 | 23 | Human polynucleoti |
| 5 | 36.4 | 15.5 | 2691 | 23 | Drosophila melanog |
| 6 | 36.4 | 15.5 | 4954 | 21 | DNA encoding a Dro |
| 7 | 36.4 | 15.5 | 8104 | 23 | Drosophila melanog |
| 8 | 33.6 | 14.3 | 4745 | 24 | Synechococcus sp. |

| | | | | | | |
|------|------|------|--------|----|-----------|--------------------|
| C 9 | 33.4 | 14.2 | 1024 | 24 | ABN74235 | Bovine embryonic g |
| C 10 | 33 | 14.0 | 1024 | 25 | ABZ82320 | Toxicologically re |
| C 11 | 33 | 14.0 | 2864 | 22 | AAH76458 | cDNA corresponding |
| C 12 | 33 | 14.0 | 2865 | 24 | ABL61798 | Colon adenocarcino |
| C 13 | 32 | 13.6 | 10587 | 22 | AAK82382 | Human immune/haema |
| C 14 | 31.6 | 13.4 | 147309 | 24 | ABK49450 | Human transporter |
| C 15 | 31.4 | 13.4 | 289 | 21 | AAA31706 | Plant microsatelli |
| C 16 | 31 | 13.2 | 10609 | 22 | AAK70162 | Human immune/haema |
| C 17 | 31 | 13.2 | 10831 | 22 | AAK70159 | Human immune/haema |
| C 18 | 31 | 13.2 | 48203 | 22 | AAK70161 | Human immune/haema |
| C 19 | 31 | 13.2 | 48203 | 22 | AAK81663 | Human immune/haema |
| C 20 | 31 | 13.2 | 48203 | 22 | AAK82628 | Human immune/haema |
| C 21 | 31 | 13.2 | 48204 | 22 | AAK70164 | Human immune/haema |
| C 22 | 31 | 13.2 | 48204 | 22 | AAK81666 | Human immune/haema |
| C 23 | 31 | 13.2 | 48204 | 22 | AAK82630 | Human immune/haema |
| C 24 | 30.8 | 13.1 | 770 | 23 | ABK50302 | Potato starch bran |
| C 25 | 30.8 | 13.1 | 3231 | 17 | AAH42632 | Class A starch bra |
| C 26 | 30.8 | 13.1 | 15004 | 22 | AAH27885 | Nucleotide sequenc |
| C 27 | 30.6 | 13.0 | 6806 | 24 | ABK97370 | Cystatin A (CSTA) |
| C 28 | 30.2 | 12.9 | 359 | 22 | AAI29247 | Colon tumour relat |
| C 29 | 30.2 | 12.9 | 359 | 24 | ABK45662 | cDNA encoding colo |
| C 30 | 30.2 | 12.9 | 359 | 25 | ABZ33433 | Human colon tumour |
| C 31 | 30.2 | 12.9 | 600 | 22 | AAK06869 | Reverse translatio |
| C 32 | 30.2 | 12.9 | 1076 | 24 | ABK76533 | Reverse translatio |
| C 33 | 30.2 | 12.9 | 1289 | 19 | AAV61202 | cDNA encoding huma |
| C 34 | 30.2 | 12.9 | 1288 | 19 | AAV48113 | Full length cDNA s |
| C 35 | 30.2 | 12.9 | 1289 | 19 | AAV61202 | Nucleotide sequenc |
| C 36 | 30.2 | 12.9 | 1289 | 19 | AAV58587 | Prostate tumour sp |
| C 37 | 30.2 | 12.9 | 1289 | 22 | AAK563558 | Human prostate cDN |
| C 38 | 30.2 | 12.9 | 1289 | 22 | AAK563558 | Human prostate tum |
| C 39 | 30.2 | 12.9 | 1289 | 22 | AAH93466 | Human prostate-spe |
| C 40 | 30.2 | 12.9 | 1289 | 22 | AAH84780 | Human prostate-spe |
| C 41 | 30.2 | 12.9 | 1289 | 22 | AAH02531 | Prostate tumour an |
| C 42 | 30.2 | 12.9 | 1289 | 22 | AAH86954 | Human P503S invent |
| C 43 | 30.2 | 12.9 | 1289 | 24 | ABK71255 | Human prostate tum |
| C 44 | 30.2 | 12.9 | 1289 | 24 | ABK58639 | Prostate tumour cD |
| C 45 | 30.2 | 12.9 | 1289 | 24 | ABL94930 | Human N1-1862 cDNA |

ALIGNMENTS

- RESULT 1
AAD44320
ID AAD44320 standard; DNA; 235 BP.
XX
AC AAD44320;
XX
DT 13-DEC-2002 (first entry)
XX
DE Spl gene fragment.
XX
KW Fibrotic condition; gene expression; cirrhosis; hypertrophic scar;
KW gene therapy; fibrosis; skin disorder; sclerodermic lesion; keloid;
KW trauma; surgery; Spl; ds.
XX
OS Unidentified.
XX
FN WO200266071-A2.
XX
PD 29-AUG-2002.
XX
PF 21-DEC-2001; 2001WO-US49141.
XX
PR 03-JAN-2001; 2001US-259585P.
XX
XX (UYJE-) UNIV JEFFERSON THOMAS.
XX
XX Mauviel A;
XX
XX WPI; 2002-667041/71.
XX
PT Treating a fibrotic condition, e.g. cirrhosis, comprises administering

12-DEC-2002.

31-MAY-2002; 2002WO-US17050.

01-JUN-2001; 2001US-295359P.

08-JUN-2001; 2001US-296878P.

08-JUN-2001; 2001US-297223P.

15-JUN-2001; 2001US-298615P.

15-JUN-2001; 2001US-298669P.

15-JUN-2001; 2001US-298693P.

21-JUN-2001; 2001US-300176P.

19-APR-2002; 2002US-373891P.

(INCV-) INCYTE GENOMICS INC.
(YUEH/) YUE H.

Yue H, Tang YT, Baughn MR, Becha SD, Warren BA, Walia NK, Lal PG;
Lee EA, Hafalia AJA, Richardson TW, Griffin JA, Emerling BM;
Ramkumar J, Yue H, Swarnakar A, Tran B, Li JX, Yao MG, Yang J;
Ison CH, Forsythe IJ, Honchell CD, Arvizu CS, Elliott VS, Lu Y;
Ding L, Luo W, Wang YE, Burford N, Borowsky ML, Nguyen DB;
Chinn AM, Kable AE;

WPI: 2003-140626/13.
P-PSDB; AAE333780.

New human nucleic acid associated proteins (NAAP), useful for
diagnosing, treating and preventing diseases or conditions associated
with the aberrant NAAP expression e.g. cancer, AIDS, atherosclerosis,
epilepsy, or infections

Claim 105; Column 247-248; 257pp; English.

The present invention relates to human nucleic acid associated proteins
(NAAP) and polynucleotides encoding such proteins. NAAP sequences are
useful in diagnosing, treating and preventing diseases or conditions
associated with the decreased expression or overexpression of NAAP such
as cell proliferative (e.g. cancer, atherosclerosis), neurological (e.g.
epilepsy, Huntington's disease, stroke), immune/inflammatory (e.g. AIDS,
allergies) and developmental (e.g. Hypothyroidism, Cushing's syndrome)
disorders or infections. They are also used in gene therapy. The present
sequence is human NAAP-20 cDNA.

Sequence 2862 BP; 753 A; 807 C; 712 G; 590 T; 0 other;

Query Match 100.0%; Score 235; DB 25; Length 2862;
Best Local Similarity 100.0%; Pred.No. 4.5e-68;
Matches 235; Conservative 0; Mismatches 0; Indels 0; Gaps 0

QY 1 ATGGGTGGCAGATCATCTCTTCCTCCCTCGGGTACCCCTACCTCAAAGGACAGAGT 60
DB 420 ATGGGTGGCAGATCATCTCTTCCTCTGGGCTACCCCTACCTCAAAGGACAGAGT 479

QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGGAGTCTTCGAAGATCCACAGTCTCTGGTG 120
DB 480 GCAGCAGTACCAATGGCAGCAATGGCAGTGGAGTCTTCGAAGATCCACAGTCTCTGGTG 539

QY 121 GSCATGATGTTGGTGGCTGGCGCTCCCAACTTACAGAACCCAGCAAGTCTCTGACAGGACTAC 180
DB 540 GSCATGATGTTGGTGGCTGGCGCTCCCAACTTACAGAACCCAGCAAGTCTCTGACAGGACTAC 599

QY 181 CTGGAGTGAATGCCATAATTCAGTATCAAGTATATCCACAGTTCACAGACCGTTGA 235
DB 600 CTGGAGTGAATGCCATAATTCAGTATCAAGTATATCCACAGTTCACAGACCGTTGA 654

RESULT 3
AAI60335
ID AAI60335 standard; cDNA; 3090 BP.
XX AAI60335;
XX AAI60335;
XX 22-OCT-2001 (first entry)

[illegible]

Best Local Similarity 54.5%; Pred. No. 0.16; Matches 73; Conservative 0; Mismatches 61; Indels 0; Gaps 0;

QY 30 GGGGCTACCCCTACCTCAAGGAACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGT 89

Db 1296 GCGGCAGCCATAGCAAGCACTGGCATCAGCAGCACTTCCATGGCAAGCAGCGCAA 1355

QY 90 GAGTCTTCCAAGATCGCACAGTCTCTGGTGGCAGTATGTTGGCTGGCTCCCAAC 149

Db 1356 TACTCAATATGCAGCAGCAACTGCCGAGGATGAGGATGTTGGATGGCGCTGCCACG 1415

QY 150 TTACAGAACCCAGCA 163

Db 1416 ATGCAGCAGCAGCA 1429

RESULT 7

ABL24298

ID ABL24298 standard; DNA; 8104 BP.

XX ABL24298;

XX 26-MAR-2002 (first entry)

DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 24367.

XX Drosophila; developmental biology; cell signalling; insecticide;

KW pharmaceutical; gene; ds.

XX Drosophila melanogaster.

OS WO200171042-A2.

PN 27-SEP-2001.

XX 23-MAR-2001; 2001WO-US09231.

XX 23-MAR-2000; 2000US-191637P.

PR 11-JUL-2000; 2000US-0614150.

XX (PEKE) PE CORP NY.

PA Venter JC, Adams M, Li FWD, Myers EW;

XX WPI; 2001-656860/75.

XX New isolated nucleic acid detection reagent for detecting 1000 or more genes from Drosophila and for elucidating cell signalling and cell-cell interactions -

PS Claim 1; SEQ ID NO 24367; 21pp + Sequence Listing; English.

XX The invention relates to an isolated nucleic acid detection reagent capable of detecting 1000 or more genes from Drosophila. The invention is useful in developmental biology and in elucidating cell signalling and cell-cell interactions in higher eukaryotes for the development of insecticides, therapeutics and pharmaceutical drugs. The invention discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA sequences (ABL01840-ABL16175) and the encoded proteins (AB57737-AB572072).

XX The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 8104 BP; 2422 A; 1863 C; 1717 G; 2102 T; 0 other;

Query Match 15.5%; Score 36.4; DB 23; Length 8104;

Best Local Similarity 54.5%; Pred. No. 0.2; Matches 73; Conservative 0; Mismatches 61; Indels 0; Gaps 0;

QY 30 GGGGCTACCCCTACCTCAAGGAACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGT 89

Db 1235 GCGGCAGCCATAGCAAGCACTTGGCATCAGCAGCACTTCCAATGGCAAGCAGCGCAA 1294

QY 90 GAGTCTTCCAAGATCGCACAGTCTCTGGTGGCAGTATGTTGGCTGGCTCCCAAC 149

Db 1295 TACTCAATATGCAGCAGCAACTGCCGAGGATGAGGATGTTGGATGGCGCTGCCACG 1354

QY 150 TTACAGAACCCAGCA 163

Db 1355 ATGCAGCAGCAGCA 1368

RESULT 8

ABS54601

ID ABS54601 standard; DNA; 4745 BP.

XX ABS54601;

XX 28-NOV-2002 (first entry)

DE Synecococcus sp. isoamylase gene.

XX ds; gene; isoamylase; PCR; gene cloning.

OS Synecococcus sp. strain pcc7942.

XX Key Location/Qualifiers

FT 1647..3731

FT /*tag= a

FT /product= "Isoamylase"

XX JP2002262877-A.

XX 17-SEP-2002.

XX 07-MAR-2001; 2001JP-0064133.

XX 07-MAR-2001; 2001JP-0064133.

XX (KAGA-) KAGAKU GIJUTSU SHINKO JIGYODAN.

PA (UYTY) UNIV TOKYO.

XX WPI; 2002-703285/76.

DR P-PSDB; ABG70853.

XX A new isoamylase, cloning a gene, a primer, an isoamylase, a gene, an organism -

XX Claim 7; Page 12-15; 22pp; Japanese.

XX The invention relates to a method for cloning a gene encoding isoamylase based on a DNA amplified by a PCR from a DNA of a genome or a library using the PCR primers appearing as ABS54602 and ABS54603. The primers may also amplify a base sequence in which part of the bases is replaced, deleted and/or added in it by a PCR. Also included are an isoamylase appearing as ABG70853 or an amino acid sequence in which part of the amino acids is deleted, replaced or added in it, a gene encoding the above isoamylase, and an organism in which the above gene is introduced to an organism in which isoamylase is deleted. The method is used for cloning a gene encoding isoamylase. The present sequence is the isoamylase gene from Synecococcus isolated using the primers of the invention.

XX Sequence 4745 BP; 930 A; 1385 C; 1258 G; 1171 T; 1 other;

Query Match 14.3%; Score 33.6; DB 24; Length 4745;

Best Local Similarity 49.2%; Pred. No. 1.4; Matches 87; Conservative 0; Mismatches 90; Indels 0; Gaps 0;

QY 56 GAGTGGCAGCAGTACCAATGGCAGCAATGGCAGTGTCTTCCAAGATGCCAGTCTC 115

Db 4251 GAGTAGCGCAGCGGTGGCGCAGCAATGCAAGTGTGTGNTACCGGGTTCATGCCCGC 4310

QY 116 TGGTGGCAGTATGTTGGCTGGCTCCCACTTACAGAACCCAGCAAGTCTTGACAG 175

Db 4311 TTGGGCTGTGCAAGAGCGGTGCCATCTCATTTTGAAGATATGCGACTGTGCGCCAG 4370

QY 176 ACTACTGGAGTGTGCTTAATATTCAGTATCCACAGTTCAGACCGT 232

Db 4371 TCTGCTCAATCTGCGCAAGAGATACTCCACAGCATTGCCCAATCCCTAAGCCCT 4427

RESULT 9

ID ABN74235/c

XX ABN74235 standard; cDNA; 1024 BP.

AC ABN74235;

XX 03-JUL-2002 (first entry)

DT DT

DE DE

XX Bovine embryonic germ (EG) cell cDNA EST #286.

XX Bovine; Bos taurus; EST; expressed sequence tag; totipotency;

KW development; gene; ss.

XX Bos taurus.

OS

XX WO200194550-A2.

PN

XX 13-DEC-2001.

PD

XX 07-JUN-2001; 2001WO-US18576.

PF

XX 07-JUN-2000; 2000US-209874P.

PR

XX 06-JUN-2001; 2001US-0876143.

PP

XX (INF1-) INF1GEN INC.

PA

XX Bilertsen KJ, Pfister-Genskow M, Childs L;

PI

XX WPI; 2002-351289/38.

DR

XX An expressed sequence tag (EST), the expression of which, or its

PT complementary sequence, in a cell identifies the cell as a

PT developmentally competent or incompetent cell.

XX

PS Example 16; Page 364; 584pp; English.

XX

XX The present invention describes an expressed sequence tag (EST), where

CC the EST is an isolated, enriched, or purified nucleic acid sequence

CC representing all or part of a gene, the expression of which, or its

CC complementary sequence, in a cell identifies the cell as a

CC developmentally competent or incompetent cell. Molecules which induce

CC developmental competence in a cell line are useful for inducing

CC totipotency in one or more cells. Molecules which induce developmental

CC competence in a cell line are useful for preventing a full term

CC pregnancy in an animal and inhibiting totipotency. The molecules are

CC also useful for treating a disease in an animal by inducing development

CC of one or more cells of the animal into a specific cell type. The

CC present sequence represents a bovine EST which is given in the

CC exemplification of the present invention.

XX

SQ Sequence 1024 BP; 187 A; 274 C; 282 G; 227 T; 54 other;

Query Match 14.2%; Score 33.4; DB 24; Length 1024;

Best Local Similarity 54.5%; Pred. No. 0.84; Mismatches 56; Indels 0; Gaps 0;

Matches 67; Conservative 0; Mismatches 56; Indels 0; Gaps 0;

QY 19 CTTCCTCTCTGGGCTACCCCTCAAGAGAGAGTGGCAGCAGTACCAATGGCA 78

Db 570 CTCCCGCTGCGGTCCACCTCGGCTCGGGGACAGGTTGCGAGCTTGAGGACGAGC 511

QY 79 GCAATGCGAGTGAAGTTCACAGATCCAGAGTCTCTGTGGCAGTATGTGTGCTG 138

Db 510 GCAAGAGGAGGAAGCCCTGGTGGATATGCCAGGCTCAGAAACGTGGCAGCTGTGAGCC 451

QY 139 CCG 141

Db 450 CCG 448

RESULT 10

ID ABZ83230/c

XX ABZ83230 standard; cDNA; 1024 BP.

AC ABZ83230;

XX 14-MAY-2003 (first entry)

DT DT

DE DE

XX Toxicologically relevant human nucleotide sequence #389.

XX Toxicologically relevant gene; toxicological response; gene; ss.

KW Homo sapiens.

XX WO2003016500-A2.

PN

XX 27-FEB-2003.

PD

XX 16-AUG-2002; 2002WO-US26514.

PF

XX 16-AUG-2001; 2001US-313080P.

PR

XX (PHAS-) PHASE-1 MOLECULAR TOXICOLOGY INC.

PA

XX Neft RE, Dunn RT, Adkins K, Pickett GG, Kier LD, Schmeisler K;

PI Alen P;

XX WPI; 2003-268322/26.

DR

XX Determining a toxicological response to an agent, useful for screening

PT of drugs, comprises comparing the expression profile of one or more

PT human toxic response genes to a reference gene expression profile

PT indicative of toxicity.

XX

PS Claim 1; Page 140; 455pp; English.

XX

XX The present invention describes a method (M1) for determining a

CC toxicological response to an agent, which comprises comparing the

CC expression profile of one or more human toxic response genes to a

CC reference gene expression profile indicative of toxicity, and so

CC determining the presence of a toxic response to the agent. Also

CC described: (1) an array comprising one or more polynucleotides selected

CC from the genes corresponding to the partial sequences given in ABZ82842

CC to ABZ84764, and their fragments of at least 20 nucleotides, or

CC homologues; and (2) determining if a gene putatively identified to be a

CC toxic response gene plays a role on toxic response pathways by

CC determining the expression profile of the gene after exposure of cells

CC or a human subject to a known toxic pharmaceutical or industrial agent,

CC comprising: (a) exposing cells to an agent or isolating cells from a

CC human subject who was exposed to an agent; (b) obtaining the test gene

CC expression profile for a putatively identified toxic response gene after

CC exposure to a known toxic pharmaceutical or industrial agent; and

CC (c) comparing the test profile to the expression profile of a gene with

CC a similar function or comparing the test profile to the expression

CC profile of that gene after exposure to other known toxic compounds. The

CC methods are useful for predicting and determining toxicological responses

CC on a cellular, organ or system level. The arrays comprising the human

CC genes are useful for toxicological screening of drugs, pharmaceutical

CC compounds and chemicals.

XX

SQ Sequence 1024 BP; 271 A; 251 C; 289 G; 213 T; 0 other;

Query Match 14.0%; Score 33; DB 25; Length 1024;

Best Local Similarity 57.1%; Pred. No. 1.1; Mismatches 60; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

Matches 60; Conservative 0; Mismatches 45; Indels 0; Gaps 0;

QY 21 TCCTCTCTCTGGGCTACCCCTCAAGAGAGAGTGGCAGCAGTACCAATGGCAGC 80

Db 134 TCCTCTCTCTGGGCTACCCCTCAAGAGAGAGTGGCAGCAGTACCCCAAGAGCAGC 75

QY 81 AATGGCAGTGTCTTCCCAAGAAATCCGACAGTCTCTGTGGGGCAG 125
 DB 74 AGGAGCAGGAAGGCTTTCCGGGGCCTCATGTAGTCGGGGCGGAG 30

RESULT 11
 AAH76458/c
 ID AAH76458 standard; cDNA; 2864 BP.

XX AAH76458;
 AC
 CC
 DE 22-OCT-2001 (first entry)
 DT
 XX cDNA corresponding to human IFN-alpha induced gene encoding ERP-70.

DE Human; interferon-alpha induced gene; type I interferon treatment;
 KW chronic viral hepatitis; relapsing remitting multiple sclerosis;
 KW neoplastic disease; IFN-alpha; interferon-alpha; ERP-70; ss.
 XX
 OS Homo sapiens.

XX Key Location/Qualifiers
 FH 46..1983
 FT /*tag= a
 FT /product= "ERP-70"
 FT

PN WO200159155-A2.

XX 16-AUG-2001.

XX 09-FEB-2001; 2001WO-GB00578.

XX 11-FEB-2000; 2000GB-0003203.

XX 11-FEB-2000; 2000GB-0003204.

PR 11-FEB-2000; 2000GB-0003205.

PR 11-FEB-2000; 2000GB-0003206.

PR 11-FEB-2000; 2000GB-0003207.

PR 11-FEB-2000; 2000GB-0003208.

PR 11-FEB-2000; 2000GB-0003210.

PR 11-FEB-2000; 2000GB-0003212.

PR 11-FEB-2000; 2000GB-0003213.

PR 11-FEB-2000; 2000GB-0003215.

PR 11-FEB-2000; 2000GB-0003220.

PR 11-FEB-2000; 2000GB-0003221.

PR 11-FEB-2000; 2000GB-0003222.

PR 17-FEB-2000; 2000GB-0003768.

XX (PHAR-) PHARMA PACIFIC PTY LTD.

XX Meritet J, Dron M, Tovey MG;

XX WPI; 2001-483570/52.

DR P-PSDB; AAG66531.

XX Predicting responsiveness of a patient to treatment with a type I
 PT interferon comprising determining the level of induced proteins after
 PT treatment with a type I interferon, -
 XX
 PS Example 1; Page 54-57; 133pp; English.

XX The invention relates to a method for predicting responsiveness of a
 CC patient to treatment with a type I interferon. The method comprises
 CC determining the level of one or more proteins with a 646, 164, 126, 598,
 CC 98, 177, 761, 361, 941, 657, 817, 429, 473, 399, 285 or 303 amino acid
 CC sequence fully defined in the specification after treatment with a
 CC type I interferon. The method allows a physician to determine whether
 CC a patient suffering from chronic viral hepatitis, neoplastic disease
 CC or relapsing remitting multiple sclerosis will respond favourably to
 CC Type I interferon treatment via oromucosal administration. The
 CC present sequence is a cDNA corresponding to an interferon-alpha
 CC induced gene that encodes one of the polypeptides listed above.
 XX
 SQ Sequence 2864 BP; 786 A; 643 C; 729 G; 706 T; 0 other;

Query Match 14.0%; Score 33; DB 22; Length 2864;
 Best Local Similarity 57.1%; Pred. No. 1.8; 45; Indels 0; Gaps 0;
 Matches 60; Conservative 0; Mismatches 0; Gaps 0;

QY 21 TCCTCCTCTGGGGCTACCCCTTACCTCAAAGAAACAGAGTGGCAGAGTACCAATGGCAGC 80
 DB 134 TCCTCGTCCGGGGCCTCGGCACCCGCCACGCGCAGAGTGCACCCGCCCAAGAGCAGC 75

QY 81 AATGGCAGTGTCTTCCCAAGAAATCCGACAGTCTCTGTGGGGCAG 125
 DB 74 AGGAGCAGGAAGGCTTTCCGGGGCCTCATGTAGTCGGGGCGGAG 30

RESULT 12

ABL61798/c

ID ABL61798 standard; DNA; 2865 BP.

XX ABL61798;

AC
 CC
 DE 15-MAY-2002 (first entry)

DT Colon adenocarcinoma related gene sequence SEQ ID NO:135.

XX Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
 KW stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
 KW cytostatic; gene therapy; antineoplastic; Wilm's tumour; adenocarcinoma;
 KW gene; ds.

XX Homo sapiens.

OS
 XX WO200194629-A2.

XX 13-DEC-2001.

XX 30-MAY-2001; 2001WO-US10838.

XX 05-JUN-2000; 2000US-209473P.

PR 05-JUN-2000; 2000US-209531P.

PR 18-SEP-2000; 2000US-233133P.

PR 18-SEP-2000; 2000US-233617P.

PR 20-SEP-2000; 2000US-234009P.

PR 20-SEP-2000; 2000US-234034P.

PR 20-SEP-2000; 2000US-234052P.

PR 22-SEP-2000; 2000US-234509P.

PR 22-SEP-2000; 2000US-234567P.

PR 25-SEP-2000; 2000US-234923P.

PR 25-SEP-2000; 2000US-234924P.

PR 25-SEP-2000; 2000US-235077P.

PR 25-SEP-2000; 2000US-235082P.

PR 25-SEP-2000; 2000US-235134P.

PR 25-SEP-2000; 2000US-235280P.

PR 26-SEP-2000; 2000US-235637P.

PR 26-SEP-2000; 2000US-235638P.

PR 27-SEP-2000; 2000US-235711P.

PR 27-SEP-2000; 2000US-235720P.

PR 27-SEP-2000; 2000US-235840P.

PR 27-SEP-2000; 2000US-235863P.

PR 28-SEP-2000; 2000US-236028P.

PR 28-SEP-2000; 2000US-236032P.

PR 28-SEP-2000; 2000US-236033P.

PR 28-SEP-2000; 2000US-236034P.

PR 28-SEP-2000; 2000US-236109P.

PR 28-SEP-2000; 2000US-236111P.

PR 29-SEP-2000; 2000US-236842P.

PR 29-SEP-2000; 2000US-236891P.

PR 02-OCT-2000; 2000US-237172P.

PR 02-OCT-2000; 2000US-237173P.

PR 02-OCT-2000; 2000US-237278P.

PR 02-OCT-2000; 2000US-237294P.

PR 02-OCT-2000; 2000US-237295P.

PR 02-OCT-2000; 2000US-237316P.

PR 03-OCT-2000; 2000US-237425P.

RESULT 13
AAK82382/C
ID AAK82382 standard: DNA: 10587 BP.

| | |
|--|--|
| RESULT 15 | |
| AAA31706/c | |
| ID AAA31706 standard; DNA; 289 BP. | |
| XX | |
| AAA31706; | |
| XX | |
| AC | |
| XX | |
| 05-JUL-2000 (first entry) | |
| XX | |
| DE | |
| Plant microsatellite marker #667. | |
| XX | |
| Plant microsatellite sequence; core repeat sequence; detection; probe; | |
| KW DNA polymorphism; genome mapping; physical mapping; fingerprinting; | |
| KW variety identification; genetic variability evaluation; primer; ss. | |
| XX | |
| Eucalyptus grandis. | |
| OS | |
| XX | |
| WO9967421-A1. | |
| XX | |
| 29-DEC-1999. | |
| XX | |
| 25-JUN-1999; 99WO-NZ00092. | |
| XX | |
| 25-JUN-1998; 98US-0105307. | |
| XX | |
| (GENE-) GENESIS RES & DEV CORP LTD & FLETCHER. | |
| PA (FLET-) FLETCHER CHALLENGE FORESTS LTD. | |
| XX | |
| Havukkala IU, Bloksberg LN, Glenn M; | |
| XX | |
| WPI; 2000-116958/10. | |
| DR | |
| XX | |
| New plant microsatellite markers and associated flanking species for | |
| PT the detection of polymorphic generic markers - | |
| XX | |
| Claim 1; Page 273; 392pp; English. | |
| PS | |

RESULT 4

US-09-030-607-111
; Sequence 111, Application US/09030607
; Patent No. 6262245
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Dillon, Davin C.
; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS FOR
; NUMBER OF SEQUENCES: 224
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED AND BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: WA
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA: US/09/030,607
; APPLICATION NUMBER: US/09/030,607
; FILING DATE: 25-FEB-1998
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Maki, David J.
; REGISTRATION NUMBER: 31,392
; REFERENCE/DOCKET NUMBER: 210121.427C3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 111:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1289 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; US-09-030-607-111

Query Match 12.9%; Score 30.2; DB 3; Length 1289;
Best Local Similarity 55.1%; Pred. No. 0.8;
Matches 59; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 17 CTCCTCTCTCTCTGGGGTACCCCTACCTCAAGGACAGAGTGGCAGCAGTACCAATGG 76
Db 852 CACTTCTGCTCTGCCACTACTGCTGCCACATGGGAACGTGAAGAGGACCCCTGGCAAG 911
QY 77 CAGCAATGGCAGTGAAGTCTTCCAAAGATCGCAGATCTCTGTGGTGGC 123
Db 912 CAGCAGTGAATGGGGAGGGGACAGGATCTTAACAATGTCACATTGGGC 958

RESULT 5

US-09-439-313-111
; Sequence 111, Application US/09439313
; Patent No. 6329505
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: Harlocker, Susan Louise
; APPLICANT: Jiang Yucui
; APPLICANT: Reed, Steven G.
; APPLICANT: Kalos, Michael
; APPLICANT: Fanger, Gary
; APPLICANT: Retter, Mark
; APPLICANT: Solk, John
; APPLICANT: Day, Craig
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THERAPY AND

; TITLE OF INVENTION: DIAGNOSIS OF PROSTATE CANCER

; FILE REFERENCE: 210121.427C9
; CURRENT APPLICATION NUMBER: US/09/439,313
; CURRENT FILING DATE: 1999-11-12
; NUMBER OF SEQ ID NOS: 575
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 111
; LENGTH: 1289
; TYPE: DNA
; ORGANISM: Homo sapien
; US-09-439-313-111

Query Match 12.9%; Score 30.2; DB 4; Length 1289;
Best Local Similarity 55.1%; Pred. No. 0.8;
Matches 59; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 17 CTCCTCTCTCTCTGGGGTACCCCTACCTCAAGGACAGAGTGGCAGCAGTACCAATGG 76
Db 852 CACTTCTGCTCTGCCACTACTGCTGCCACATGGGAACGTGAAGAGGACCCCTGGCAAG 911
QY 77 CAGCAATGGCAGTGAAGTCTTCCAAAGATCGCAGATCTCTGTGGTGGC 123
Db 912 CAGCAGTGAATGGGGAGGGGACAGGATCTTAACAATGTCACATTGGGC 958

RESULT 6

US-09-352-616A-111
; Sequence 111, Application US/09352616A
; Patent No. 6395278
; GENERAL INFORMATION:
; APPLICANT: Dillon, Davin C.
; APPLICANT: Harlocker, Susan Louise
; APPLICANT: Jiang, Yucui
; APPLICANT: Xu, Jiangchun
; APPLICANT: Mitcham, Jennifer Lynn
; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY AND DIAGNOSIS
; FILE REFERENCE: 210121.427C8
; CURRENT APPLICATION NUMBER: US/09/352,616A
; CURRENT FILING DATE: 1999-07-13
; NUMBER OF SEQ ID NOS: 472
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 111
; LENGTH: 1289
; TYPE: DNA
; ORGANISM: Homo sapien
; US-09-352-616A-111

Query Match 12.9%; Score 30.2; DB 4; Length 1289;
Best Local Similarity 55.1%; Pred. No. 0.8;
Matches 59; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 17 CTCCTCTCTCTCTGGGGTACCCCTACCTCAAGGACAGAGTGGCAGCAGTACCAATGG 76
Db 852 CACTTCTGCTCTGCCACTACTGCTGCCACATGGGAACGTGAAGAGGACCCCTGGCAAG 911
QY 77 CAGCAATGGCAGTGAAGTCTTCCAAAGATCGCAGATCTCTGTGGTGGC 123
Db 912 CAGCAGTGAATGGGGAGGGGACAGGATCTTAACAATGTCACATTGGGC 958

RESULT 7

US-09-232-149A-111
; Sequence 111, Application US/09232149A
; Patent No. 6465611
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer Lynn
; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE
; FILE REFERENCE: 210121.427C6
; CURRENT APPLICATION NUMBER: US/09/232,149A

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; CURRENT FILING DATE: 1999-01-15
; NUMBER OF SEQ ID NOS: 338
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 111
; LENGTH: 1289
; TYPE: DNA
; ORGANISM: Homo sapien
US-09-232-149A-111

Query Match          12.9%; Score 30.2; DB 4; Length 1289;
Best Local Similarity 55.1%; Pred. No. 0.8;
Matches 59; Conservative 0; Mismatches 48; Indels 0; Gaps 0;

QY 17 CTCCTCCCTCTGGGTACCCCTACTCTCAAGACAGAGTGGCAGCAGTACCAATGG 76
DB 852 CACTCTGCTCTGCCACTACTGTGTCACATGGGAATGTGAAGAGCGCACCTTGGAAG 911
QY 77 CAGCAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGTGGGC 123
DB 912 CAGCAGTGTGGGGAGGGGACAGGATCTAACATGTCACTTGGGC 958

RESULT 8
US-09-328-352-2777/c
; Sequence 1, Application US/09328352
; Patent No. 6562958
; GENERAL INFORMATION:
; APPLICANT: Gary L. Brston et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO ACINETOBACTER
; FILE REFERENCE: GT099-03PA
; CURRENT APPLICATION NUMBER: US/09/328,352
; CURRENT FILING DATE: 1999-06-04
; NUMBER OF SEQ ID NOS: 8252
; SEQ ID NO 2777
; LENGTH: 621
; TYPE: DNA
; ORGANISM: Acinetobacter baumannii
US-09-328-352-2777

Query Match          12.6%; Score 29.6; DB 4; Length 621;
Best Local Similarity 68.3%; Pred. No. 0.86;
Matches 41; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

QY 68 TACCAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGTGGCAGTA 127
DB 202 TAAGATGTAGCAGAGGAATGAGTATCTACGCATCACACCATCACTGCTGTCTACTA 143

RESULT 9
US-08-793-273C-1
; Sequence 1, Application US/08793273C
; Patent No. 6482410
; GENERAL INFORMATION:
; APPLICANT: Crossin, Kathryn L.
; APPLICANT: Phillips, Greg
; APPLICANT: Prieto, Anne L.
; TITLE OF INVENTION: CYTOTACTIN DERIVATIVES THAT STIMULATE ATTACHMENT AND
; FILE REFERENCE: BEC00228
; FILE REFERENCE: BEC00228
; CURRENT APPLICATION NUMBER: US/08/793,273C
; CURRENT FILING DATE: 1997-05-22
; PRIOR APPLICATION NUMBER: PCT/US95/11684
; PRIOR FILING DATE: 1995-09-14
; PRIOR APPLICATION NUMBER: 08/308,359
; PRIOR FILING DATE: 1994-09-16
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 7286
; TYPE: DNA
; ORGANISM: Homo Sapiens
; FEATURE:

; NAME/KEY: CDS
; LOCATION: (55)...(6654)
US-08-793-273C-1

Query Match          12.4%; Score 29.2; DB 4; Length 7286;
Best Local Similarity 50.0%; Pred. No. 4.6;
Matches 73; Conservative 0; Mismatches 73; Indels 0; Gaps 0;

QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGTG 120
DB 2789 GGAGGAATGGCAAGCAGCTATTGACAGTTACAGAAATTAAGTATGCCCCCACTCTGTGAG 2848
QY 121 GCAGTATGTTGTGCTCGCGCTCCCACTTACAGAACCCAGCAAGTTCTGACAGGACTAC 180
DB 2849 GGAGCAGCTAGTGTGATGTTCCAAAGAGCCACAGCCACCAACCAAAACCACTCA 2908
QY 181 CTGGAGTGTGCTTAATATTCAGTAT 206
DB 2909 CAGGTCTGAGGCGGGAATGAATAT 2934

RESULT 10
PCT-US95-11684-1
; Sequence 1, Application PC/TUS9511684
; GENERAL INFORMATION:
; APPLICANT: THE SCRIPPS RESEARCH INSTITUTE
; TITLE OF INVENTION: CYTOTACTIN DERIVATIVES THAT STIMULATE
; TITLE OF INVENTION: ATTACHMENT AND NEURITE OUTGROWTH, AND METHODS OF MAKING
; NUMBER OF INVENTION: AND USING SAME
; NUMBER OF SEQUENCES: 28
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: The Scripps Research Institute, Office of
; ADDRESSER: Patent Counsel
; STREET: 10666 North Torrey Pines Road, TPC 8
; CITY: La Jolla
; STATE: CA
; COUNTRY: USA
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC Compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/11684
; FILING DATE: 14-SEP-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/308,359
; FILING DATE: 16-SEP-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Logan, April C.
; REGISTRATION NUMBER: 33,950
; REFERENCE/DOCKET NUMBER: BEC0019P
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-554-2937
; TELEFAX: 619-554-6312
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7286 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 55..6654
; OTHER INFORMATION: /product= "cytotactin"
PCT-US95-11684-1

Query Match          12.4%; Score 29.2; DB 5; Length 7286;
Best Local Similarity 50.0%; Pred. No. 4.6;
Matches 73; Conservative 0; Mismatches 73; Indels 0; Gaps 0;
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QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGGCTTCCAGAAATCCACAGTCTCTGGTG 120
Db 2789 GGAGGAATGGCAAGCGCAGCTATTGACAGATTACAGAAATTAAGTATGCCGCCCATCTCTGGAG 2848
QY 121 GGCAGTATGTTGGCTGGCGCTCCCACTTACAGAACCAAGTCTTCGACAGGACTAC 180
Db 2849 GGCACACGCTGAGTTGATGTTCCAAAGAGCCACAGCCACCAACCAACCACTCA 2908
QY 181 CTGAGTATGCTTATATTCAGTAT 206
Db 2909 CAGGCTGAGCGCGGAAGTGAATAT 2934

RESULT 11

US-08-958-201-1/c
; Sequence 1, Application US/08958201
; Patent No. 5977319
; GENERAL INFORMATION:
; APPLICANT: Pope, Anthony R
; APPLICANT: Pritchard, Kevin
; APPLICANT: Williams, Andrew J
; APPLICANT: Johnson, Kevin S
; TITLE OF INVENTION: Specific binding members for estradiol;
; TITLE OF INVENTION: materials and methods
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall O'Toole Gerstein Murray & Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606-6402
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25 (BPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/958,201
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/028,897
; FILING DATE: 21-OCT-1996
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 354 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; CLONE: D12
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..354
; US-08-958-201-1

Query Match 12.3%; Score 29; DB 2; Length 354;
Best Local Similarity 54.1%; Pred. No. 1.1;
Matches 59; Conservative 0; Mismatches 50; Indels 0; Gaps 0;
QY 78 AGCAATGGCAGTCTCCAGAAATCGCAGTCTCTGGTGGCAGTATGTTGGCT 137
Db 318 ATCAAAACCATTTAGTACTCCTAAATCGCAGTAAATATATGGCGGTCTCTCGGCTG 259
QY 138 GCGCTCCCAACTTACAGAACCAAGCAAGTCTTCAGAGGACTACCTGGAG 186
Db 258 CAGGCTGTTCTTCAGATACACCGCTGTTCTTGGATTGTCCCTGGAG 210

RESULT 12

US-09-831-206-1/c
; Sequence 1, Application US/09831206

; Patent No. 6573070
; GENERAL INFORMATION:
; APPLICANT: MacNeil, Douglas J.
; APPLICANT: Weinberg, David H.
; APPLICANT: Van der Ploeg, Leonardus H. T.
; TITLE OF INVENTION: DNA MOLECULES ENCODING THE MELANOCORTIN
; TITLE OF INVENTION: 4 RECEPTOR PROTEIN FROM RHESUS MONKEY
; FILE REFERENCE: 20190P
; CURRENT APPLICATION NUMBER: US/09/831,206
; CURRENT FILING DATE: 2001-06-28
; PRIOR APPLICATION NUMBER: PCT/US99/25767
; PRIOR FILING DATE: 1999-11-05
; PRIOR APPLICATION NUMBER: 60/107,721
; PRIOR FILING DATE: 1998-11-09
; NUMBER OF SEQ ID NOS: 6
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 1030
; TYPE: DNA
; ORGANISM: rhesus monkey (Macaca mulatta)
; US-09-831-206-1

Query Match 12.3%; Score 28.8; DB 4; Length 1030;
Best Local Similarity 65.6%; Pred. No. 2.2;
Matches 42; Conservative 0; Mismatches 22; Indels 0; Gaps 0;
QY 93 TCTTCCAGATCGCACAGTCTCTGGTGGCAGTATGTTGGCTGCCGCTCCCACTTA 152
Db 115 TTTTCCAGGAGCACTGCTGGCATTGCTGTCAGTCTGTCGCTGCCAGAGGTG 56
QY 153 CAGA 156
Db 55 CAGA 52

RESULT 13

US-09-238-303-7
; Sequence 7, Application US/09238303B
; Patent No. 6284253
; GENERAL INFORMATION:
; APPLICANT: Barr, Margaret C.
; TITLE OF INVENTION: No. 6284253el Feline Immunodeficiency Virus Nucleotide Sequence
; FILE REFERENCE: 18617.0059
; CURRENT APPLICATION NUMBER: US/09/238,303B
; CURRENT FILING DATE: 1999-01-28
; EARLIER APPLICATION NUMBER: US 60/072,927
; EARLIER FILING DATE: 1998-01-29
; NUMBER OF SEQ ID NOS: 17
; SEQ ID NO 7
; LENGTH: 9751
; TYPE: DNA
; ORGANISM: Unknown
; FEATURE:
; OTHER INFORMATION: recombinant viral clone constructed from the genomic DNA of
; OTHER INFORMATION: a Pallas's cat feline immunodeficiency virus
; US-09-238-303-7

Query Match 12.3%; Score 28.8; DB 3; Length 9751;
Best Local Similarity 56.2%; Pred. No. 7.4;
Matches 54; Conservative 0; Mismatches 42; Indels 0; Gaps 0;
QY 121 GGCAGTATGTTGGCTGGCGCTCCCACTTACAGAACCAAGTCTTCAGAGGACTAC 180
Db 2045 GGCAGAGGGGAGAGCTGCTGCCCTATCAAAAGTGCAGCAATTTCAAAACAGCAT 2104
QY 181 CTGGAGTGATGCTTAATATTCAGTATCAAGTATCC 216
Db 2105 CAACACTCAGATCAGCAACATGTCATATATAC 2140

RESULT 14

US-09-946-239-7
; Sequence 7, Application US/09946239

Patent No. 6579527
GENERAL INFORMATION:
APPLICANT: Barz, Margaret C.
TITLE OF INVENTION: No. 6579527el Feline Immunodeficiency Virus Nucleotide and
FILE OF INVENTION: Polypeptide Sequences
FILE REFERENCE: 18617.0059
CURRENT APPLICATION NUMBER: US/09/946.239
CURRENT FILING DATE: 2001-09-04
PRIOR APPLICATION NUMBER: US 09/238.303, US 60/072,927
PRIOR FILING DATE: 1999-01-28, 1998-01-29
NUMBER OF SEQ ID NOS: 17
SEQ ID NO 7
LENGTH: 9751
TYPE: DNA
ORGANISM: Unknown
FEATURE:
OTHER INFORMATION: recombinant viral clone constructed from the genomic DNA of
OTHER INFORMATION: a Pallas's cat feline immunodeficiency virus
US-09-946-239-7

Query Match 12.3%; Score 28.8; DB 4; Length 9751;
Best Local Similarity 56.2%; Pred. No. 7.4;
Matches 54; Conservative 0; Mismatches 42; Indels 0; Gaps 0;
QY 121 GCAGTATGTTGTGGCTGCCGCTCCCACTTACAGAACCGACGAGTCTGTGACGAGACTAC 180
DB 2045 GCGAGAGGGGAGAGCTGCTGCCCTATCAACCAAGTGCAGCAATTTCAACACGAGTAT 2104
QY 181 CTGGAGTGTGCTAATATTCAGTATCAAGTAATCC 216
DB 2105 CAACACTCAGATCAGCAACATGTCATTAATAC 2140

RESULT 15
US-09-634-238-409
Sequence 409, Application US/09634238
Patent No. 6544772
GENERAL INFORMATION:
APPLICANT: Glenn, Matthew
APPLICANT: Havukala, Ilkka J.
APPLICANT: Bloksberg, Leonard, N.
APPLICANT: Lubbers, Mark W.
APPLICANT: Dekker, James
APPLICANT: Christensen, Anna C.
APPLICANT: Holland, Ross
APPLICANT: O'Toole, Paul W.
APPLICANT: Reid, Julian R.
APPLICANT: Coolbear, Timothy
TITLE OF INVENTION: Polynucleotides, materials incorporating
FILE OF INVENTION: them and methods for using them.
FILE REFERENCE: 11000.1043U1
CURRENT APPLICATION NUMBER: US/09/634,238
CURRENT FILING DATE: 2000-08-08
NUMBER OF SEQ ID NOS: 422
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 409
LENGTH: 3269
TYPE: DNA
ORGANISM: Lactobacillus rhamnosus
US-09-634-238-409

Query Match 12.1%; Score 28.4; DB 4; Length 3269;
Best Local Similarity 50.7%; Pred. No. 5.7;
Matches 68; Conservative 0; Mismatches 66; Indels 0; Gaps 0;
QY 20 TTCCTCTCTGGGCTACCCCTACTCAAGAAACAGAGTGGCAGCTACCAATGGCAG 79
DB 1343 TCCCTTCGCTCAAGTGGGCTACTTCAAGTGAAGTGGTAATCAGCAATTCATCACATT 1402
QY 80 CAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGTGGGAGATGTTGTGGCTGC 139
DB 1403 GGCCTCAACATCATCGAGCATGTGGGGACCGCTCCGTTGGGTAACCTGTACCGCTGC 1462

QY 140 CGCTCCCACTTAC 153
DB 1463 CTCATGGACTTAC 1476
Search completed: February 18, 2004, 16:10:41
Job time : 63.179 secs

Db 121 GGCAGTATGTTGTGGCTCCCGCTCCCACTTACAGAACCGACAGTCTTGACAGACTAC 180
Qy 181 CTGGAGTATGCTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235
Db 181 CTGGAGTATGCTTATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235

RESULT 2

US-10-117-722-436
; Sequence 436, Application US/10117722
; Publication No. US20030219744A1
; GENERAL INFORMATION:
; APPLICANT: Tang, Y. Tom
; APPLICANT: Liu, Chenghua
; APPLICANT: Asundi, Vinod
; APPLICANT: Zhang, Jie
; APPLICANT: Drmanac, Radoje T.
; TITLE OF INVENTION: Polypeptides
; FILE REFERENCE: 784CIP2B
; CURRENT APPLICATION NUMBER: US/10/117,722
; PRIOR FILING DATE: 2002-04-04
; PRIOR APPLICATION NUMBER: 09/620,312
; PRIOR FILING DATE: 2000-07-19
; PRIOR APPLICATION NUMBER: 09/552,317
; PRIOR FILING DATE: 2000-04-25
; PRIOR APPLICATION NUMBER: 09/488,725
; PRIOR FILING DATE: 2000-01-21
; NUMBER OF SEQ ID NOS: 1104
; SOFTWARE: pt_FL_genes Version 1.0
; SEQ ID NO 436
; LENGTH: 3289
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (201)...(2558)
US-10-117-722-436

Query Match 100.0%; Score 235; DB 13; Length 3289;
Best Local Similarity 100.0%; Pred. No. 1e-71;
Matches 235; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
Db 514 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 573
Qy 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAATCCACAGTCTCTGGTG 120
Db 574 GCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAATCCACAGTCTCTGGTG 633
Qy 121 GCAGTATGTTGGTGGCGCTCCCACTTACAGAACCGACAGTCTTGACAGACTAC 180
Db 634 GCAGTATGTTGGTGGCGCTCCCACTTACAGAACCGACAGTCTTGACAGACTAC 693
Qy 181 CTGGAGTATGCTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235
Db 694 CTGGAGTATGCTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 748

RESULT 3

US-10-037-270-436
; Sequence 436, Application US/10037270
; Publication No. US20030104529A1
; GENERAL INFORMATION:
; APPLICANT: Tang, Y. Tom
; APPLICANT: Liu, Chenghua
; APPLICANT: Asundi, Vinod
; APPLICANT: Zhang, Jie
; APPLICANT: Ren, Feiyan
; APPLICANT: Chen, Rui-hong
; APPLICANT: Zhao, Qing A.

; APPLICANT: Wehrman, Tom
; APPLICANT: Xue, Aidong J.
; APPLICANT: Yang, Yonghong
; APPLICANT: Wang, Jian-Rui
; APPLICANT: Zhou, Ping
; APPLICANT: Ma, Yunging
; APPLICANT: Wang, Dunrui
; APPLICANT: Wang, Zhiwei
; APPLICANT: Tillinghast, John
; APPLICANT: Drmanac, Radoje T.
; TITLE OF INVENTION: No. US20030104529A1el Nucleic Acids and
; TITLE OF INVENTION: Polypeptides
; FILE REFERENCE: 784CIP2B
; CURRENT APPLICATION NUMBER: US/10/037,270
; CURRENT FILING DATE: 2002-01-04
; PRIOR APPLICATION NUMBER: 09/552,317
; PRIOR FILING DATE: 2000-04-25
; PRIOR APPLICATION NUMBER: 09/488,725
; PRIOR FILING DATE: 2000-01-21
; NUMBER OF SEQ ID NOS: 1104
; SOFTWARE: pt_FL_genes Version 1.0
; SEQ ID NO 436
; LENGTH: 3289
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (201)...(2558)
US-10-037-270-436

Query Match 100.0%; Score 235; DB 15; Length 3289;
Best Local Similarity 100.0%; Pred. No. 1e-71;
Matches 235; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
Db 514 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 573
Qy 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAATCCACAGTCTCTGGTG 120
Db 574 GCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAAATCCACAGTCTCTGGTG 633
Qy 121 GCAGTATGTTGGTGGCGCTCCCACTTACAGAACCGACAGTCTTGACAGACTAC 180
Db 634 GCAGTATGTTGGTGGCGCTCCCACTTACAGAACCGACAGTCTTGACAGACTAC 693
Qy 181 CTGGAGTATGCTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235
Db 694 CTGGAGTATGCTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 748

RESULT 4

US-09-814-353-18291
; Sequence 18291, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Lee, John
; APPLICANT: Thompson, Pamela
; APPLICANT: Lillie, James
; TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
; TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND
; TITLE OF INVENTION: THERAPY OF OVARIAN CANCER
; FILE REFERENCE: MRI-006B
; CURRENT APPLICATION NUMBER: US/09/814,353
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: US 60/191,031
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: US 60/207,124
; PRIOR FILING DATE: 2000-05-25
; PRIOR APPLICATION NUMBER: US 60/211,940
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: US 60/216,820
; PRIOR FILING DATE: 2000-07-07

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; PRIOR APPLICATION NUMBER: US 60/220,661
; PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
; NUMBER OF SEQ ID NOS: 22037
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 18291
; LENGTH: 570
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-814-353-18291

Query Match      29.7%; Score 69.8; DB 13; Length 570;
Best Local Similarity 91.4%; Pred. No. 5.3e-14;
Matches 74; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

2y 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGACAGAGTG 60
   |||||
Db 430 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGACAGAGTG 489
   |||||

2y 61 GCAGCAGTACCAATGGCAGCA 81
   |||||
Db 490 GCAGCAGTACCTCGGCGGCCA 510
   |||||

RESULT 5
; US-09-814-353-21704
; Sequence 21704, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Lee, John
; APPLICANT: Thompson, Pamela
; APPLICANT: Lillie, James
; TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
; TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND
; FILE REFERENCE: MRI-006B
; CURRENT APPLICATION NUMBER: US/09/814,353
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: US 60/191,031
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: US 60/207,124
; PRIOR FILING DATE: 2000-05-25
; PRIOR APPLICATION NUMBER: US 60/211,940
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: US 60/216,820
; PRIOR FILING DATE: 2000-07-07
; PRIOR APPLICATION NUMBER: US 60/220,661
; PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
; NUMBER OF SEQ ID NOS: 22037
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 21704
; LENGTH: 682
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 682
; OTHER INFORMATION: n = A,T,C or G
; US-09-814-353-21704

Query Match      27.8%; Score 65.4; DB 13; Length 682;
Best Local Similarity 98.5%; Pred. No. 2e-12;
Matches 66; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

2y 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGACAGAGTG 60
   |||||
Db 615 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGACAGAGTG 674
   |||||

2y 61 GCAGCAG 67
   |||||
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Db 675 GCAGCAG 681

RESULT 6
; US-09-814-353-5620
; Sequence 5620, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Lee, John
; APPLICANT: Thompson, Pamela
; APPLICANT: Lillie, James
; TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
; TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND
; FILE REFERENCE: MRI-006B
; CURRENT APPLICATION NUMBER: US/09/814,353
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: US 60/191,031
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: US 60/207,124
; PRIOR FILING DATE: 2000-05-25
; PRIOR APPLICATION NUMBER: US 60/211,940
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: US 60/216,820
; PRIOR FILING DATE: 2000-07-07
; PRIOR APPLICATION NUMBER: US 60/220,661
; PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
; NUMBER OF SEQ ID NOS: 22037
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5620
; LENGTH: 411
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-814-353-5620

Query Match      21.3%; Score 50; DB 13; Length 411;
Best Local Similarity 100.0%; Pred. No. 4.2e-07;
Matches 50; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAG 50
   |||||
Db 362 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAG 411
   |||||

RESULT 7
; US-09-814-353-11907
; Sequence 11907, Application US/09814353
; Publication No. US20030165831A1
; GENERAL INFORMATION:
; APPLICANT: Lee, John
; APPLICANT: Thompson, Pamela
; APPLICANT: Lillie, James
; TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR
; TITLE OF INVENTION: IDENTIFICATION, ASSESSMENT, PREVENTION, AND
; FILE REFERENCE: MRI-006B
; CURRENT APPLICATION NUMBER: US/09/814,353
; CURRENT FILING DATE: 2001-03-21
; PRIOR APPLICATION NUMBER: US 60/191,031
; PRIOR FILING DATE: 2000-03-21
; PRIOR APPLICATION NUMBER: US 60/207,124
; PRIOR FILING DATE: 2000-05-25
; PRIOR APPLICATION NUMBER: US 60/211,940
; PRIOR FILING DATE: 2000-06-15
; PRIOR APPLICATION NUMBER: US 60/216,820
; PRIOR FILING DATE: 2000-07-07
; PRIOR APPLICATION NUMBER: US 60/220,661
; PRIOR FILING DATE: 2000-07-25
; PRIOR APPLICATION NUMBER: US 60/257,672
; PRIOR FILING DATE: 2000-12-21
; NUMBER OF SEQ ID NOS: 22037
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; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 147309
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(147309)
; OTHER INFORMATION: n = A,T,C or G
US-10-436-185-3

Query Match      13.4%; Score 31.6; DB 13; Length 147309;
Best Local Similarity 58.5%; Pred. No. 14;
Matches 55; Conservative 0; Mismatches 39; Indels 0; Gaps 0;

QY 130 TTGTGGTGGCGCTCCCAACTTACAGAACCCAGCAAGTCTTGACAGGACTACTCTGGAGTCA 189
DB 97837 TTGTGCTGCGCCCTCCACCCCATGACTCAGTCTCTTCAATGCGAGCTAGAGTGA 97896

QY 190 TGCTTAATATTCAGTATCAAGTAAATCCACAGTT 223
DB 97897 TCCATTTAAACGTAATGTAGTCTTCTCATTTGT 97930

RESULT 12
US-10-085-117-74
; Sequence 74, Application US/10085117
; Publication No. US20030232334A1
; GENERAL INFORMATION:
; APPLICANT: Morris, David W.
; APPLICANT: Engelhard, Eric K.
; TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR CANCER
; FILE REFERENCE: 529452000121
; CURRENT APPLICATION NUMBER: US/10/085,117
; CURRENT FILING DATE: 2002-02-27
; PRIOR APPLICATION NUMBER: US 09/798,586
; PRIOR FILING DATE: 2001-03-02
; NUMBER OF SEQ ID NOS: 361
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 74
; LENGTH: 4452
; TYPE: DNA
; ORGANISM: Mus musculus
US-10-085-117-74

Query Match      13.3%; Score 31.2; DB 12; Length 4452;
Best Local Similarity 58.7%; Pred. No. 4.5;
Matches 54; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 45 TCAAGGACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAT 104
DB 2842 TGATAGGAATGTGTGTCCACCTTTCCAAAGAAAGAAAGACAAAGAGATTTCAGATT 2901

QY 105 CGCACAGTCTCTGGTGGGCGAGTATGTTGGC 136
DB 2902 GGCCAGCTCTGATAGTGAATTTTITGCC 2933

RESULT 13
US-10-085-117-73/c
; Sequence 73, Application US/10085117
; Publication No. US20030232334A1
; GENERAL INFORMATION:
; APPLICANT: Morris, David W.
; APPLICANT: Engelhard, Eric K.
; TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR CANCER
; FILE REFERENCE: 529452000121
; CURRENT APPLICATION NUMBER: US/10/085,117
; CURRENT FILING DATE: 2002-02-27
; PRIOR APPLICATION NUMBER: US 09/798,586
; PRIOR FILING DATE: 2001-03-02
; NUMBER OF SEQ ID NOS: 361
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; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 73
; LENGTH: 99934
; TYPE: DNA
; ORGANISM: Mus musculus
; FEATURE:
; NAME/KEY: variation
; LOCATION: (1)..(99934)
; OTHER INFORMATION: n = any nucleotide
US-10-085-117-73

Query Match      13.3%; Score 31.2; DB 12; Length 99934;
Best Local Similarity 58.7%; Pred. No. 16;
Matches 54; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 45 TCAAGGACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAT 104
DB 11611 TGATAGGAATGTGTGTCCACCTTTCCAAAGAAAGAAAGACAAAGAGATTTCAGATT 11552

QY 105 CGCACAGTCTCTGGTGGGCGAGTATGTTGGC 136
DB 11551 GGCCAGCTCTGATAGTGAATTTTITGCC 11520

RESULT 14
US-10-056-454A-18
; Sequence 18, Application US/10056454A
; Publication No. US20030166919A1
; GENERAL INFORMATION:
; APPLICANT: National Starch and Chemical Investment Holding Corporation
; TITLE OF INVENTION: Improvements in or Relating to Plant Starch Composition
; NUMBER OF SEQUENCES: 20
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: National Starch and Chemical Investment Holding Corporation
; STREET: 1000 Uniqema Blvd.
; CITY: Newcastle
; STATE: Delaware
; COUNTRY: United States of America
; ZIP: 19720
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/10/056,454A
; FILING DATE: 25-Jun-2002
; INFORMATION FOR SEQ ID NO: 18:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3231 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 18:
US-10-056-454A-18

Query Match      13.1%; Score 30.8; DB 13; Length 3231;
Best Local Similarity 54.4%; Pred. No. 5.4;
Matches 62; Conservative 0; Mismatches 52; Indels 0; Gaps 0;

QY 41 TACCTCAAAGAACACAGAGTGGCAGCAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAA 100
DB 536 TTCAACAATGGACACAGCAGTACCAATGAACTGAGAACGATGAGTTGAGCGGTCAAG 595

QY 101 GAATGCGACAGTCTCTGGTGGGCGAGTATGTTGGTGGCGCTCCCAACTTACA 154
DB 596 TGATCTTACAGGAAGTGTGAGAGAGTTGGATTTTGTTCATCACTACAACATA 649

RESULT 15
US-10-199-676-23/c
; Sequence 23, Application US/10199676
; Publication No. US20040014051A1
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; GENERAL INFORMATION:
; APPLICANT: Vickie L. Brown-Driver
; APPLICANT: Kenneth W. Dobie
; TITLE OF INVENTION: ANTISENSE MODULATION OF BREAST CANCER-1 EXPRESSION
; FILE REFERENCE: PTS-0017
; CURRENT APPLICATION NUMBER: US/10/199,676
; CURRENT FILING DATE: 2002-07-18
; NUMBER OF SEQ ID NOS: 84
; SEQ ID NO 23
; LENGTH: 130001
; TYPE: DNA
; ORGANISM: H. sapiens
; FEATURE:
US-10-199-676-23

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Query Match      13.1%; Score 30.8; DB 12; Length 130001;
Best Local Similarity 48.8%; Pred. No. 25;
Matches 83; Conservative 0; Mismatches 87; Indels 0; Gaps 0;

QY      3  GGCTGGCAGATCATCTCTTCTCTCTCTGGGGCTACCCCTACCTCAAAGGAACAGAGTGGC 62
Db      128564  GGCAGGATGGTCTCGAACTCTGACCTCGTGATCCGGCTGCTCAGCCTCCCAAGTGCT 128605

QY      63  AGCAGTACCAATGGCAGCAATGGCAGTGAGTCTTCCAAAGAAATGCACAGTCTCTGTGGG 122
Db      128604  GGGATTACAGGCGTGAGCCACCGTGCCCGCAGCATGGCTAATTTTGTAGAGACAGGCTTT 128545

QY      123  CAGTATGTTGGTGGCTGGCGCTCCCACTTACAGAACCCAGCAAGTCTTGAC 172
Db      128544  CAGCATGTTGGCCAGGCTGGTCTCGAACTCTTGACCTCACATGATCTGCC 128495

```

Search completed: February 18, 2004, 16:15:54
Job time : 277.233 secs

GenCore version 5.1.6
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DN nucleic - nucleic search, using sw model

run on: February 18, 2004, 12:26:35 ; Search time 2369.2 Seconds
(without alignments)
2410.749 Million cell updates/sec

Title: US-10-026-341a-1

Perfect score: 235

Sequence: 1 atgggtgcagatcatctct.....ccacagttccagaccgttga 235

Scoring table:

IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:

1: em_estba.*

2: em_esthum.*

3: em_estin.*

4: em_estmu.*

5: em_estov.*

6: em_estpl.*

7: em_estro.*

8: em_hic.*

9: gb_estl.*

10: gb_est2.*

11: gb_hic.*

12: gb_est3.*

13: gb_est4.*

14: gb_est5.*

15: em_estfun.*

16: em_estom.*

17: em_gss_hum.*

18: em_gss_inv.*

19: em_gss_pln.*

20: em_gss_vrt.*

21: em_gss_fun.*

22: em_gss_mam.*

23: em_gss_mus.*

24: em_gss_pro.*

25: em_gss_rod.*

26: em_gss_phg.*

27: em_gss_vrl.*

28: gb_gssl.*

29: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----|--------------------|
| 1 | 235 | 100.0 | 755 | 12 | BI559366 603253037 |
| 2 | 233.4 | 99.3 | 645 | 9 | AW609227 RC3-ST018 |
| 3 | 233.4 | 99.3 | 949 | 13 | BQ890167 AGENCOURT |
| 4 | 222.4 | 94.6 | 1150 | 13 | BQ072120 AGENCOURT |

| | | | | | |
|----|-------|------|------|----|----------|
| 5 | 192.6 | 82.0 | 546 | 12 | BG803579 |
| 6 | 192.6 | 82.0 | 617 | 9 | AW822630 |
| 7 | 186.8 | 79.5 | 590 | 10 | BE937236 |
| 8 | 183.2 | 78.0 | 653 | 14 | BY738990 |
| 9 | 183 | 77.9 | 586 | 13 | BU702181 |
| 10 | 157.4 | 67.0 | 416 | 9 | AW445976 |
| 11 | 147.2 | 62.6 | 404 | 14 | CB811553 |
| 12 | 75.4 | 32.1 | 681 | 12 | BM427106 |
| 13 | 70 | 29.8 | 504 | 12 | BG997913 |
| 14 | 67.2 | 28.6 | 880 | 13 | BU906803 |
| 15 | 67.2 | 28.6 | 928 | 14 | CA787807 |
| 16 | 61.8 | 26.3 | 922 | 14 | CA987845 |
| 17 | 61.4 | 26.1 | 670 | 9 | AL647036 |
| 18 | 61.4 | 26.1 | 671 | 13 | BQ397449 |
| 19 | 58 | 24.7 | 450 | 10 | BF875151 |
| 20 | 57 | 24.3 | 382 | 10 | BF510251 |
| 21 | 50.4 | 21.4 | 629 | 14 | CD279637 |
| 22 | 50.4 | 21.4 | 755 | 14 | CA470150 |
| 23 | 50.4 | 21.4 | 832 | 14 | CA476365 |
| 24 | 48.8 | 20.8 | 473 | 14 | CD280695 |
| 25 | 48.8 | 20.8 | 902 | 14 | CA469702 |
| 26 | 48.4 | 20.6 | 740 | 14 | CB940752 |
| 27 | 40 | 17.0 | 257 | 9 | AW178522 |
| 28 | 39.8 | 16.9 | 502 | 10 | BE696452 |
| 29 | 39.4 | 16.8 | 622 | 14 | CD282914 |
| 30 | 38.4 | 16.3 | 954 | 13 | BU236388 |
| 31 | 36.2 | 15.4 | 810 | 12 | BI837393 |
| 32 | 35.8 | 15.2 | 527 | 14 | CD036456 |
| 33 | 35.8 | 15.2 | 642 | 14 | CA622180 |
| 34 | 35.6 | 15.1 | 744 | 10 | BG480269 |
| 35 | 34.6 | 14.7 | 675 | 9 | AW027138 |
| 36 | 34.6 | 14.7 | 804 | 12 | BI837784 |
| 37 | 34.6 | 14.7 | 1201 | 9 | AL523072 |
| 38 | 34.6 | 14.7 | 1201 | 9 | AL529564 |
| 39 | 34.4 | 14.6 | 874 | 14 | CD302448 |
| 40 | 34.2 | 14.6 | 1201 | 9 | AL523073 |
| 41 | 33.8 | 14.4 | 448 | 29 | BZ867528 |
| 42 | 33.8 | 14.4 | 673 | 29 | AG096615 |
| 43 | 33.8 | 14.4 | 1201 | 13 | BX394709 |
| 44 | 33.6 | 14.3 | 273 | 9 | AV328003 |
| 45 | 33.4 | 14.2 | 859 | 14 | CB559451 |

ALIGNMENTS

RESULT 1
BI559366
LOCUS 603253037F1 NIH_MGC_97 Homo sapiens cDNA clone IMAGE:5295465 5', linear EST 05-SEP-2001
DEFINITION mRNA sequence.
ACCESSION BI559366
VERSION BI559366.1 GI:15446680
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 755)
NIH-MGC <http://mgc.ncbi.nlm.nih.gov/>.
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE Unpublished
JOURNAL
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs@mail.nih.gov
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
CDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki Toshiyuki and Piero Carninci (RIKEN)
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Plate: LLAM11747 row: j column: 10

FEATURES
source
High quality sequence stop: 754.
Location/Qualifiers
1. .755

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5295465"
/lab_host="DH10B"
/clone_lib="NIH_MGC_97"
/note="Organ: testis; Vector: pBluescriptR (modified pBluescript KS+); Site 1: BamHI; Site 2: SalI-XhoI (gtcgag
); Oligo-dr primed using primer 5'-TTTTTTTTTTTTTTTNN-3',
size-selected for average insert size 2.2 kb and
normalized to ROT 5. This is a primary library enriched
for full-length clones and constructed using the
Cap-trapper method (Carninci, in preparation). Library
constructed by M. Brownstein (NIMH/NHGRI, National
Institutes of Health). Note: this is a NIH_MGC Library."
199 a 203 c 207 g 146 t

BASE COUNT
ORIGIN
Query Match 100.0%; Score 235; DB 12; Length 755;
Best Local Similarity 100.0%; Pred. No. 8.7e-58;
Matches 235; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGTACCCCTTACCTCAAGGAACACAGAGTG 60
DB 353 ATGGCTGGCAGATCATCTCTCTCTCTGGGGTACCCCTTACCTCAAGGAACACAGAGTG 412
QY 61 GCAGCAGTACCAATGGCAGCAGTGTCTTCCAGAAATCGCAGTCTCTGGTG 120
DB 413 GCAGCAGTACCAATGGCAGCAGTGTCTTCCAGAAATCGCAGTCTCTGGTG 472
QY 121 GGCAGTATGTTGTGGCTGGCTGCCCTCCCACTTACAGAACCCAGTGTCTGACAGACTAC 180
DB 473 GGCAGTATGTTGTGGCTGGCTGCCCTCCCACTTACAGAACCCAGTGTCTGACAGACTAC 532
QY 181 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235
DB 533 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 587

RESULT 2
AW609227
LOCUS
DEFINITION
RC3-ST0186-300100-017-f03 ST0186 Homo sapiens cDNA, mRNA sequence.
ACCESSION
AW609227
VERSION
AW609227.1 GI:7313968
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 645)
ECGP <http://www.ludwig.org.br/ORESTES>.
The FAPESP/LICR Human Cancer Genome Project
Unpublished
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(<http://www.ludwig.org.br/scripts/gethtml2.pl?l1=RC3&t2=RC3-ST0186-300100-017-f03&t3=2000-01-30&t4=1>)
Seq primer: puc 18 forward
High quality sequence stop: 644.
Location/Qualifiers
1. .645

FEATURES
source
High quality sequence stop: 754.
Location/Qualifiers
1. .755

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="ST0186"
/note="Organ: stomach; Vector: puc18; Site 1: SmaI;
Site 2: SmaI; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the pUC 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."
158 a 152 c 189 g 146 t

BASE COUNT
ORIGIN
Query Match 99.3%; Score 233.4; DB 9; Length 645;
Best Local Similarity 99.6%; Pred. No. 2.4e-57;
Matches 234; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGTACCCCTTACCTCAAGGAACACAGAGTG 60
DB 334 ATGGCTGGCAGATCATCTCTCTCTCTGGGGTACCCCTTACCTCAAGGAACACAGAGTG 393
QY 61 GCAGCAGTACCAATGGCAGCAGTGTCTTCCAGAAATCGCAGTCTCTGGTG 120
DB 394 GCAGCAGTACCAATGGCAGCAGTGTCTTCCAGAAATCGCAGTCTCTGGTG 453
QY 121 GGCAGTATGTTGTGGCTGGCTGCCCTCCCACTTACAGAACCCAGTGTCTGACAGACTAC 180
DB 454 GGCAGTATGTTGTGGCTGGCTGCCCTCCCACTTACAGAACCCAGTGTCTGACAGACTAC 513
QY 181 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 235
DB 514 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCACAGTTCACAGACCGTTGA 568

RESULT 3
BQ890167
LOCUS
DEFINITION
AGENCOURT_8730196 NIH_MGC_47 Homo sapiens cDNA clone IMAGE:6339589
5', mRNA sequence.
ACCESSION
BQ890167
VERSION
BQ890167.1 GI:22282181
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 949)
NIH-MGC <http://mgc.nci.nih.gov/>.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgabbs-r@mail.nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Rubin Laboratory
DNA Sequencing by: The I.M.A.G.E. Consortium (LLNL)
Cloned by: Agencourt Bioscience Corporation
Cloned through the I.M.A.G.E. Consortium information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LLCM2534 row: k column: 14
High quality sequence stop: 570.
Location/Qualifiers
1. .949

FEATURES
source
High quality sequence stop: 570.
Location/Qualifiers
1. .949

FEATURES
source
High quality sequence stop: 644.
Location/Qualifiers
1. .645

EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GCACGAG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NIH_MGC Library."

BASE COUNT 239 a 269 c 258 g 193 t
ORIGIN

Query Match 99.3%; Score 233.4; DB 13; Length 949;
Best Local Similarity 99.6%; Pred. No. 2.9e-57;
Matches 234; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATGGGTGGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGACAGAGTG 60
DB 411 ATGGGTGGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGACAGAGTG 470

QY 61 GCAGCAGTACCACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGGTG 120
DB 471 GCAGCAGTACCACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGGTG 530

QY 121 GGCAGTATGTTGTGGCTGCCCTCCAACTTACAGAACCGAAGTCTTGACAGGACTAC 180
DB 531 GGCAGTATGTTGTGGCTGCCCTCCAACTTACAGAACCGAAGTCTTGACAGGACTAC 590

QY 181 CTGGAGTATGCTTCAATATTCAGTATCAAGTAATCCACAGATTCACAGCCGTTGA 235
DB 591 CTGGAGTATGCTTCAATATTCAGTATCAAGTAATCCACAGATTCACAGCCGTTGA 645

RESULT 4
BQ072120
LOCUS BQ072120.1 1150 bp mRNA linear EST 02-APR-2002
DEFINITION AGENCOURT 6859825 NIH_MGC_47 Homo sapiens cDNA clone IMAGE:5928633
5' mRNA sequence.

ACCESSION BQ072120
VERSION BQ072120.1 GI:19901166
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 1150)
AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: ATCC

cDNA Library Preparation: Rubin Laboratory
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LICM2105 row: h column: 10
High quality sequence stop: 571.
Location/Qualifiers

FEATURES
source
1..1150
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:5928633"
/tissue_type="neuroblastoma, cell line"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH_MGC_47"
/note="Organ: brain; Vector: pOTB7; Site 1: XhoI; Site 2: EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GCACGAG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in

the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NIH_MGC Library."

BASE COUNT 281 a 344 c 290 g 235 t
ORIGIN

Query Match 94.6%; Score 222.4; DB 13; Length 1150;
Best Local Similarity 99.2%; Pred. No. 5.2e-54;
Matches 234; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 1 ATGGGTGGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGACAGAGTG 60
DB 411 ATGGGTGGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGACAGAGTG 470

QY 61 GCAGCAGTACCACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGGTG 120
DB 471 GCAGCAGTACCACCAATGGCAGCAATGGCAGTGTCTTCCAGAAATCGCACAGTCTCTGGTG 530

QY 121 GGCAGTATGTTGTGGCTGCCCTCCAACTTACAGAACCGAAGTCTTGACAGGACTAC 180
DB 531 GGCAGTATGTTGTGGCTGCCCTCCAACTTACAGAACCGAAGTCTTGACAGGACTAC 590

QY 181 CTGG-AGTATGCTTCAATATTCAGTATCAAGTAATCCACAGATTCACAGCCGTTGA 235
DB 591 CTGGAGTATGCTTCAATATTCAGTATCAAGTAATCCACAGATTCACAGCCGTTGA 646

RESULT 5
BG803579
LOCUS BG803579
DEFINITION 0233-17 Mouse E14.5 retina lambda ZAP II Library Mus musculus cDNA, mRNA sequence.

ACCESSION BG803579.1 GI:17950485
VERSION BG803579
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 546)
AUTHORS Mu, X., Zhao, S., Pershad, R., Hsieh, T.-F., Scarpa, A., Wang, S.W., White, R.A., Beremand, P.D., Thomas, T.D., Gan, L. and Klein, W.H.
TITLE Gene expression in the developing mouse retina by EST sequencing and microarray analysis
JOURNAL Nucleic Acids Res. 29 (24), 4983-4993 (2001)
MEDLINE 21671825
PUBMED 11812828
COMMENT Contact: Klein WH
Department of Biochemistry and Molecular Biology
University of Texas M.D. Anderson Cancer Center
Box 117, 1515 Holcombe Blvd., Houston, TX 77030, USA
Tel: 713 792 3646
Fax: 713 790 0329.

FEATURES
source
1..546
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/tissue_type="neural retina"
/dev_stage="embryonic day 14.5 post-fertilization"
/clone_lib="Mouse E14.5 retina lambda ZAP II Library"
Location/Qualifiers

BASE COUNT 139 a 154 c 129 g 124 t
ORIGIN

Query Match 82.0%; Score 192.6; DB 12; Length 546;
Best Local Similarity 92.3%; Pred. No. 1.7e-45;
Matches 217; Conservative 0; Mismatches 9; Indels 9; Gaps 1;

QY 1 ATGGCTGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGACAGAGTG 60
DB 18 ATGGCTGCAGATCATCTCTTCTCTCTCTGGGGTACCCCTACCTCAAGGACAGAGTG 77

Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, T., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Koya, S., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N., Saito, R., Sakazume, N., Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Takeda, Y., Waki, K., Watahiki, A., Muramatsu, M. and Hayashizaki, Y. Direct Submission

Computational Analysis of Full-length Mouse cDNAs Compared with Human Genome Sequences *Mamm. Genome*. 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. *Genome Res.* 10 (10), 1617-1630 (2000)

RIKEN Integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. *Genome Res.* 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Please visit our web site (<http://genome.asc.riken.go.jp>) for

```

Location/Qualifiers
1. .653
/organism="Mus musculus"
/mol_time="10min"

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/clone="I920028B19"
/sex="female"
/tissue_type="amnion"

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/dev.stages="17 days pregnant adult"
/clone.lib="RIKEN full-length enriched, 17 days pregnant
adult female amnion"
149 a 199 c 180 g 122 t 3 others

78.0%; Score 183.2; DB 14; Length 653;
Similarity 89.4%; Pred. No.1e-42;
110; Conservative 0; Mismatches 16; Indels 9; Gaps 1;

1 1 ATGGCTGGCAGATCATCTCTTCTCCCTCTGGGGCTACCCCTACCTCAAAGGAACAGAGTG 60
3 ATGGCTGGCAGATCATCTCTTCTCCCTCTGGGGCTACCCCTACCTCANAGGAACAGAGTG 462

1 GCACAGTACCAATGGCAGCAATGGCAGTGGTCTTCCAGAATCGCACAGTCTCTGGTG 120
3 GGCAGATTACCAATGGCAGC-----GAGTCTTCCAGAACCGCACAGTCTCTGGTG 513

1 GGCAGTAGTGTGGCTGCGGCTCCCAACTTACAGAACCCAGCAAGTCTTGACAGCACTAC 180
4 GGCAGTAGTGTGGCTGCTACCCCAACTTACAGAACCAAGTCTTGACAGGCTCTCC 573

1 CTGGAGTGATGCGCTTAATTTCAGTATCAAGTAATCCCAAGTTCACAGACCGTTGA 235

4 CTGGAGTAATGCCTAATATTCAGTATCAAGTAATCCACAGNTCCAGACTATTGA 628

BU702181 586 bp mRNA linear EST 09-OCT-2002
 UI-MF10-by-r-a-11-0-UI.r1 NIH EMAP_F10 Mus musculus CDNA clone
 IMAGE: 5698066 5', mRNA sequence.
 BU702181
 BU702181.1 GI:23626729
 EST.
 Mus musculus (house mouse)
 Mus musculus
 Mus musculus
 Eukaryota: Metazoa: Chordata: Vertebrata: Euteleostomi:

REFERENCE 1 (bases 1 to 586)
 AUTHORS NIH-MGC http://mgs.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabbs-r@mail.nih.gov
 Tissue Procurement: Dr. Jim Lin, University of Iowa
 cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
 cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 This clone was contributed by the Brain Molecular Anatomy Project (BMAP)

Seq primer: pYX-5.
 Location/Qualifiers
 1..586
 /organism="Mus musculus"
 /mol_type="mRNA"
 /strain="C57BL/6"
 /db_xref="taxon:10090"
 /clone="IMAGE: 5638066"
 /tissue_type="whole brain"
 /dev_stage="embryo 12.5dpc"
 /lab_host="NIH BMAP F10"
 /clone_lib="NIH BMAP F10"
 /note="Organ: Brain; Vector: pYX-Asc; Site 1: EcoR I; Site 2: Not I; The library was constructed according to Bonaïdo, Lennon and Soares, Genome Research, 6:791-806, 1996. Denatured RNA was size fractionated on a 1% agarose gel. First strand cDNA synthesis was primed with oligo-dT primer containing a Not I site. Double strand cDNA was size selected according to mRNA size fraction, ligated with EcoR I adaptor, digested with NotI and then cloned directionally into pYX-Asc vector. The library tag sequence located between the Not I site and the polyA tail is CAGCAGGAC. This library was created for the University of Iowa Brain Anatomy Project (BMAP). Gene Discovery in the Developing Mouse Nervous System", supported by National Institute of Mental Health (NIMH), Hemin Chin, Ph.D., program coordinator."

BASE COUNT 143 a 167 c 161 g 115 t

Query Match 77.9%; Score 183; DB 13; Length 586;
 Best Local Similarity 89.8%; Pred. No. 1.1e-42;
 Matches 211; Conservative 0; Mismatches 15; Indels 9; Gaps 1;
 QY 1 ATGGCTGGCAGATCATCTCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 60
 Db 196 ATGGCTGGCAGATCATCTCTCTCTGGGGCTACCCCTACCTCAAGGAACAGAGTG 255
 QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGAGTCTTCCAAAGTAATCGCAGTCTCTGGTG 120
 Db 256 GGAACAGTACCAATGGGAGC-----GAGCTCTCCAAAGAACCCGACAGTCTCTGGTG 306
 QY 121 GCAGATGTTGTGGCTGCGCTGCCAACTTACAGAACCCAGCAAGTCTTGACGAGCTAC 180
 Db 307 GGCAGTGTGTGTGGCTGTCTACCCCACTTACAGAACCCAGCAAGTCTTGACGAGTCTCC 366
 QY 181 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCCAAGTTCAGACCGTTGA 235
 Db 367 CTGGAGTAATGCTACTATTTCAGTATCAAGTAATCCCAAGTTCAGACCTGTTGA 421

RESULT 10
 AW445976
 LOCUS 83265 MARC 1B0V Bos taurus cDNA 5', mRNA linear EST 25-APR-2001
 DEFINITION AW445976
 ACCESSION AW445976
 VERSION AW445976.1 GI:6987761

KEYWORDS
 SOURCE
 ORGANISM

Bos taurus (cow)
 Bos taurus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovidae; Bovinae; Bos.

REFERENCE
 AUTHORS

Smith, T.P.L., Grose, W.M., Freking, B.A., Roberts, A.J., Stone, R.T., Casas, E., Wray, J.E., White, J., Cho, J., Fahrenkrug, S.C., Bennett, G.L., Heaton, M.P., Laegreid, W.M., Rohrer, G.A., Chitko-McKown, C.G., Pertea, G., Hol, I., Karamycheva, S., Liang, F., Quackenbush, J. and Keele, J.W.

Sequence evaluation of four pooled-tissue normalized bovine cDNA libraries and construction of a gene index for cattle Genome Res. 11 (4), 626-630 (2001)

JOURNAL
 MEDLINE
 PUBMED

11282978

Contact: Smith TPL
 USDA, ARS, US Meat Animal Research Center
 PO Box 166, Clay Center, NE 68933-0166, USA
 Tel: 402 762 4366
 Fax: 402 762 4390

Email: smith@email.marc.usda.gov
 Single pass sequencing. Bases called and trimmed with phred v0.980904.e. Vector identified by cross_match with the -minscore 20 and -minmatch 12 options.

PCR Primers

FORWARD: AGGAAACAGCTATGACCAT

BACKWARD: GTTTCCAGTCACGACG

Plate: 45 row: B column: 16

Seq primer: ATTTAGTGCACCTATAG.

FEATURES
 source

Location/Qualifiers

1..416

/organism="Bos taurus"

/mol_type="mRNA"

/db_xref="taxon:9913"

/tissue_type="pooled"

/lab_host="DH10B"

/clone_lib="MARC 1B0V"

/notes="Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI; Library made from pooled tissue from lymph node, ovary, fat, hypothalamus, and pituitary."

BASE COUNT 113 a 114 c 104 g 85 t

Query Match 67.0%; Score 157.4; DB 9; Length 416;
 Best Local Similarity 93.7%; Pred. No. 2.8e-35;
 Matches 164; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

QY 61 GCAGCAGTACCAATGGCAGCAATGGCAGTGAGTCTTCCAAAGTAATCGCAGTCTCTGGTG 120

Db 1 GCAGCAGTACCAACGGGACAAATGGCAGTGAGTCTTCCAAAGTAATCGCAGTCTCGGTG 60

QY 121 GCAGTATGTTGGTGCGGCTCCCAACTTACAGAACCCAGCAAGTCTTGACGAGCTAC 180

Db 61 GCAGTATGTTGGTGCGGCTCCCAACTTACAGAACCCAGCAAGTCTTGACGAGCTAC 120

QY 181 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCCAAGTTCAGACCGTTGA 235

Db 121 CTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCCAAGTTCAGACCGTTGA 175

RESULT 11
 CB811553

LOCUS

CB811553 404 bp mRNA linear EST 16-MAY-2003
 AMGNNUC:SRPB2-00245-E5-A srpb2 (10220) Rattus norvegicus cDNA clone srpb2-00245-e5 5', mRNA sequence.

DEFINITION

CB811553

VERSION

CB811553.1 GI:29934519

KEYWORDS

Rattus norvegicus (Norway rat)

SOURCE

Rattus norvegicus

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

High quality sequence start: 8
High quality sequence stop: 504.

FEATURES

source
1. 504
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone_lib="HRI298"
/dev_stage="Adult"
/note="Organ: head neck; Vector: puc18; Site 1: SmaI;
Site 2: SmaI; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the pUC 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."

BASE COUNT 113 a 115 c 148 g 127 t 1 others
ORIGIN

Query Match 29.8%; Score 70; DB 12; Length 504;
Best Local Similarity 93.6%; Pred. No. 9.6e-10;
Matches 73; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 158 CCAGCAAGTTCGACAGGACTACCTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCC 217
DB 494 CCCAGAACCTCTGACAGGACTACCTGGAGTGATGCTTAATATTCAGTATCAAGTAATCCC 435
QY 218 ACAGTTCAGACCGTTGA 235
DB 434 ACAGTTCAGACCGTTGA 417

RESULT 14

BU906803
LOCUS
DEFINITION
IMAGE:6631848 5', mRNA linear EST 17-OCT-2002
AGENCOURT_10456317 NICHHD_XGC_001 Xenopus laevis cDNA clone
IMAGE:6631848 5', mRNA sequence.

ACCESSION BU906803

VERSION BU906803.1 GI:24088717

KEYWORDS

SOURCE

ORGANISM
Xenopus laevis (African clawed frog)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Amphibia; Batrachia; Anura; Mesobatrachia; Pipiloidea; Pipidae;
Xenopodinae; Xenopus.

REFERENCE 1 (bases 1 to 880)

AUTHORS
TITLE
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

JOURNAL

COMMENT

Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
Tissue Procurement: Martha Rebert, Steven L. Klein, Ph.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA sequencing by: Agencourt Bioscience Corporation
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LLAM14200 row: d column: 24
High quality sequence stop: 647.

FEATURES

source
1. 880
/organism="Xenopus laevis"
/mol_type="mRNA"
/db_xref="taxon:8355"
/clone="IMAGE:6631848"
/tissue_type="embryo (stage 10)"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NICHHD_XGC_Emb1"

/note="Vector: pCMV-SPORT6; Site 1: NotI; Site 2: SalI;
Cloned unidirectionally. Primer: Oligo dt. Average insert
size 1.55 kb. Constructed by Life Technologies. Note: This

is a Xenopus Gene Collection (XGC) library."

BASE COUNT 238 a 243 c 190 g 207 t 2 others
ORIGIN

Query Match 28.6%; Score 67.2; DB 13; Length 880;
Best Local Similarity 69.7%; Pred. No. 8.4e-09;
Matches 106; Conservative 0; Mismatches 43; Indels 3; Gaps 1;

QY 84 GGAGTGAGTCTTCCAGAAATCGACAGTCTCTGGTGGCAGTATGTGTGGCTCCGCT 143
DB 300 GGAGATGCTTCTTCTTAAACCGCCATAGCCCTGGGCACT---TTGTGTAGTGTGA 356
QY 144 CCCAACTTACAGAACAGCAAGTCTTGACAGCACTACCTGGAGTGATGCTTAATATTCAG 203
DB 357 CCAGTGTGCAAAATCAGAGGTTTGACCACTTTCAGAGTGTGATGCCCAATTCAG 416
QY 204 TATCAAGTAATCCAGATTCAGACCGGTGA 235
DB 417 TACCAAGTCATACCAATTCAGACTGTGA 448

RESULT 15

CA787807

LOCUS

DEFINITION

IMAGE:6859147 5', mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Xenopus laevis (African clawed frog)

Xenopus laevis

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Amphibia; Batrachia; Anura; Mesobatrachia; Pipiloidea; Pipidae;

Xenopodinae; Xenopus.

REFERENCE 1 (bases 1 to 928)

AUTHORS

TITLE

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index

JOURNAL

COMMENT

Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
Tissue Procurement: Martha Rebert, Steven L. Klein, Ph.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA sequencing by: Agencourt Bioscience Corporation
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LLAM14471 row: g column: 18
High quality sequence stop: 744.

FEATURES

source

1. 928
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/db_xref="taxon:8355"
/clone="IMAGE:6859147"
/tissue_type="oocytes"
/lab_host="DH10B (phage-resistant)"
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/note="Vector: pCMV-SPORT6; Site 1: NotI; Site 2: SalI;
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BASE COUNT 251 a 249 c 201 g 223 t 4 others
ORIGIN

Query Match 28.6%; Score 67.2; DB 14; Length 928;
Best Local Similarity 69.7%; Pred. No. 8.7e-09;
Matches 106; Conservative 0; Mismatches 43; Indels 3; Gaps 1;

QY 84 GGAGTGAGTCTTCCAGAAATCGACAGTCTCTGGTGGCAGTATGTGTGGCTCCGCT 143
DB 329 GGAGATGCTTCTTCTTAAACCGCCCAATAGCCCTGGGCACT---TTGTGTAGTGTGA 395

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model
Run on: February 18, 2004, 15:26:16 ; Search time 1542 Seconds
(without alignments)
583.865 Million cell updates/sec

Title: US-10-026-341A-2
Perfect score: 22
Sequence: 1 attcgatcgggggggggcgagc 22

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2888711 seqs, 2045481386 residues
Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : GenEmbl:

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- 2: gb_hgt:*
- 3: gb_in:*
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- 30: em_htg_hum:*
- 31: em_htg_inv:*
- 32: em_htg_other:*
- 33: em_htg_mus:*
- 34: em_htg_pln:*
- 35: em_htg_rdt:*
- 36: em_htg_mam:*
- 37: em_htg_vrt:*
- 38: em_sy:*
- 39: em_htgo_hum:*
- 40: em_htgo_mus:*
- 41: em_htgo_other:*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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| 2 | 22 | 100.0 | 22 | 6 | AR070792 | Sequence |
| 3 | 22 | 100.0 | 22 | 6 | AX195274 | Sequence |
| 4 | 22 | 100.0 | 22 | 6 | AX526281 | Sequence |
| 5 | 22 | 100.0 | 22 | 6 | BD105476 | Prolifera |
| 6 | 22 | 100.0 | 48 | 6 | AX377574 | Sequence |
| 7 | 22 | 100.0 | 48 | 6 | AX377574 | Sequence |
| 8 | 22 | 100.0 | 48 | 6 | AX127457 | Sequence |
| 9 | 22 | 100.0 | 49 | 6 | AX127457 | Sequence |
| 10 | 22 | 100.0 | 49 | 6 | AX127459 | Sequence |
| 11 | 22 | 100.0 | 49 | 6 | AX127459 | Sequence |
| 12 | 22 | 100.0 | 49 | 6 | AX377573 | Sequence |
| 13 | 21 | 95.5 | 21 | 6 | AX104633 | Sequence |
| 14 | 21 | 95.5 | 21 | 6 | AX104634 | Sequence |
| 15 | 21 | 95.5 | 21 | 6 | AX355087 | Sequence |
| 16 | 21 | 95.5 | 21 | 6 | AX355201 | Sequence |
| 17 | 21 | 95.5 | 21 | 6 | AX547686 | Sequence |
| 18 | 21 | 95.5 | 21 | 6 | AX547687 | Sequence |
| 19 | 20 | 90.9 | 20 | 6 | A89791 | Sequence 13 |
| 20 | 20 | 90.9 | 20 | 6 | A90878 | Sequence 13 |
| 21 | 20 | 90.9 | 20 | 6 | AX023404 | Sequence |
| 22 | 20 | 90.9 | 20 | 6 | AX455627 | Sequence |
| 23 | 20 | 90.9 | 20 | 6 | BD056782 | Pharmaceu |
| 24 | 19 | 86.4 | 46 | 6 | I72381 | Sequence 12 |
| 25 | 19 | 86.4 | 46 | 6 | I72382 | Sequence 13 |
| 26 | 18.8 | 85.5 | 49 | 6 | AX377573 | Sequence |
| 27 | 18.4 | 83.6 | 22 | 6 | E07877 | Synthetic n |
| 28 | 18 | 81.8 | 46 | 6 | I72381 | Sequence 12 |
| 29 | 18 | 81.8 | 46 | 6 | I72382 | Sequence 13 |
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| 31 | 17.8 | 80.9 | 274015 | 2 | AC127795 | Gasterost |
| 32 | 17.8 | 80.9 | 301660 | 2 | AC112802 | Rattus no |
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| 34 | 17.4 | 79.1 | 1123 | 9 | HSA243936 | Homo sapi |
| 35 | 17.4 | 79.1 | 7125 | 1 | AF335479 | Agrobacte |
| 36 | 17.4 | 79.1 | 9646 | 6 | AX346590 | Sequence |
| 37 | 17.4 | 79.1 | 30377 | 9 | HSLUCA1 | Human DNA s |
| 38 | 17.4 | 79.1 | 106278 | 8 | AC132215 | Rattus no |
| 39 | 17.4 | 79.1 | 126323 | 8 | AC132215 | Genomic s |
| 40 | 17.4 | 79.1 | 14969 | 8 | AC137547 | Oryza sat |
| 41 | 17.4 | 79.1 | 158266 | 2 | AC092556 | Oryza sat |
| 42 | 17.4 | 79.1 | 184461 | 2 | AC118476 | Mus muscu |
| 43 | 17.4 | 79.1 | 192324 | 9 | AC096920 | Homo sapi |
| 44 | 17.4 | 79.1 | 201964 | 10 | MMHQC29N7 | Mus muscu |
| 45 | 17.4 | 79.1 | 201986 | 10 | AC006289 | Mus muscu |

ALIGNMENTS

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| RESULT 1 | AR060640 | Sequence 12 from patent US 5840832. | 22 bp | DNA | linear | PAT 29-SEP-1999 |
| LOCUS | AR060640 | Sequence 12 from patent US 5840832. | | | | |
| DEFINITION | AR060640 | Sequence 12 from patent US 5840832. | | | | |
| ACCESSION | AR060640 | Sequence 12 from patent US 5840832. | | | | |
| VERSION | AR060640.1 | GI:5987090 | | | | |
| KEYWORDS | AR060640.1 | GI:5987090 | | | | |
| SOURCE | Unknown. | | | | | |
| ORGANISM | Unknown. | | | | | |
| REFERENCE | Unclassified. | | | | | |
| AUTHORS | 1 (bases 1 to 22) | | | | | |
| TITLE | Ono,S.Jeremy. and Strominger,J.L. | | | | | |
| JOURNAL | Transcription factor regulating MHC expression, CDNA and genomic clones encoding same and retroviral expression constructs thereof | | | | | |
| | Patent: US 5840832-A 12 24-NOV-1998; | | | | | |

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FEATURES                               Location/Qualifiers
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Qy 1 ATTTCGATCGGGCGGGCGGCGAGC 22
Db 1 ATTTCGATCGGGCGGGCGGCGAGC 22

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DEFINITION Sequence 12 from patent US 5908762.
ACCESSION  AR070792
VERSION     AR070792.1 GI:7221680
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unknown.
REFERENCE   1 (bases 1 to 22)
AUTHORS    Ono,S.Jeremy. and Strominger,J.L.
TITLE      Transcription factor regulating XMC expression CDNA and genomic
            clones encoding same and retroviral expression constructs thereof
JOURNAL    Patent: US 5908762-A 12 01-JUN-1999;
FEATURES   Location/Qualifiers
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DEFINITION Sequence 10 from Patent WO0151671.
ACCESSION  AX195274
VERSION     AX195274.1 GI:15385825
KEYWORDS
SOURCE      synthetic construct
ORGANISM    synthetic construct
            artificial sequences.
REFERENCE   1
AUTHORS     McCarthy,J. and Cordell,B.
TITLE      Methods for identifying inhibitors of neuronal degeneration
JOURNAL    Patent: WO 0151671-A 10 19-JUL-2001;
            Scios Inc. (US)
FEATURES   Location/Qualifiers
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RESULT 4
LOCUS      AX526281      22 bp      DNA      linear      PAT 21-NOV-2002
DEFINITION Sequence 2 from Patent WO02066071.
ACCESSION  AX526281
VERSION     AX526281.1 GI:25171091
KEYWORDS
SOURCE      unidentified
ORGANISM    unidentified
            unclassified.
REFERENCE   1
AUTHORS     Mauviel,A.
TITLE      Blocking spi transcription factor broadly inhibits extracellular
            matrix gene expression in vitro and in vivo: implications for the
            treatments of tissue fibrosis
JOURNAL    Patent: WO 02066071-A 2 29-AUG-2002;
            Thomas Jefferson University (US)
FEATURES   Location/Qualifiers
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RESULT 5
LOCUS      BD105476      22 bp      DNA      linear      PAT 27-AUG-2002
DEFINITION Proliferating agents and apoptosis-suppressing agents for
            pancreatic beta-cells of the islets of langerhans, and screening of
            candidate compounds thereof.
ACCESSION  BD105476
VERSION     BD105476.1 GI:22651050
KEYWORDS    WO 0193899-A/9.
SOURCE      synthetic construct
ORGANISM    synthetic construct
            artificial sequences.
REFERENCE   1 (bases 1 to 22)
AUTHORS     Okamoto,H.
TITLE      Proliferating agents and apoptosis-suppressing agents for
            pancreatic beta-cells of the islets of langerhans, and screening of
            candidate compounds thereof
JOURNAL    Patent: WO 0193899-A 9 13-DEC-2001;
            HIROSHI OKAMOTO
COMMENT     OS Artificial Sequence
            PN WO 0193899-A/9
            PD 13-DEC-2001
            PF 01-JUN-2001 WO 2001JP004660
            PR 02-JUN-2000 JP 00P 170447,28-FEB-2001 JP 01P 54072 PI
            PC A61K38/20,A61K31/573,A61K38/22,A61K48/00,A61K45/00,A61K31/455,
            PC A61K31/165,
            PC A61P43/00,A61P3/10,G01N33/15,G01N33/50,C12Q1/02//C12N15/12, PC
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            CC Description of Artificial Sequence:an artificially synthesized
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            FH Key Location/Qualifiers
            FT source 1..22

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FEATURES
source

Location/Qualifiers

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QY 1 ATTCGATCGGGCGGGCGGAGC 22

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RESULT 6

AX377574

LOCUS

Sequence 51 from Patent WO0212553.

ACCESSION AX377574

VERSION AX377574.1 GI:19573760

KEYWORDS

SOURCE

ORGANISM

synthetic construct

artificial sequences.

REFERENCE 1

AUTHORS Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrensdoerf,H. and

Muth,J.

TITLE Method for detecting mutations in nucleotide sequences

JOURNAL Patent: WO 0212553-A 51 14-FEB-2002;

Nanogen Recognomics GmbH (DE)

FEATURES

source

Location/Qualifiers

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Hairpin-Oligonucleotid"

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Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22

Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 7

AX377574/c

LOCUS

Sequence 51 from Patent WO0212553.

ACCESSION AX377574

VERSION AX377574.1 GI:19573760

KEYWORDS

SOURCE

ORGANISM

synthetic construct

artificial sequences.

REFERENCE 1

AUTHORS Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrensdoerf,H. and

Muth,J.

TITLE Method for detecting mutations in nucleotide sequences

JOURNAL Patent: WO 0212553-A 51 14-FEB-2002;

Nanogen Recognomics GmbH (DE)

FEATURES

source

Location/Qualifiers

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Best Local Similarity 100.0%; Pred. No. 45;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 8

AX127457

LOCUS

Sequence 1 from Patent WO0131057.

ACCESSION AX127457

VERSION AX127457.1 GI:14134020

KEYWORDS

SOURCE

ORGANISM

synthetic construct

artificial sequences.

REFERENCE 1

AUTHORS Muth,J. and Windhab,N.

TITLE Double-strand nucleic acid probes and the use thereof

JOURNAL Patent: WO 0131057-A 1 03-MAY-2001;

Aventis Research & Technologies GmbH & Co KG (DE)

FEATURES

source

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/organism="synthetic construct"

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Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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AX127457/c

LOCUS

Sequence 1 from Patent WO0131057.

ACCESSION AX127457

VERSION AX127457.1 GI:14134020

KEYWORDS

SOURCE

ORGANISM

synthetic construct

artificial sequences.

REFERENCE 1

AUTHORS Muth,J. and Windhab,N.

TITLE Double-strand nucleic acid probes and the use thereof

JOURNAL Patent: WO 0131057-A 1 03-MAY-2001;

Aventis Research & Technologies GmbH & Co KG (DE)

FEATURES

source

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RESULT 10
AX127459
LOCUS AX127459 49 bp DNA linear PAT 15-MAY-2001
DEFINITION Sequence 3 from Patent WO0131057.
ACCESSION AX127459
VERSION AX127459.1 GI:14134028
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1 Muth, J. and Windhab, N.
AUTHORS Double-strand nucleic acid probes and the use thereof
TITLE Patent: WO 0131057-A 3 03-MAY-2001
JOURNAL Aventis Research & Technologies GmbH & Co KG (DE)
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BASE COUNT 6 a 16 c 16 g 11 t
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Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 11
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LOCUS AX127459 49 bp DNA linear PAT 15-MAY-2001
DEFINITION Sequence 3 from Patent WO0131057.
ACCESSION AX127459
VERSION AX127459.1 GI:14134028
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1 Muth, J. and Windhab, N.
AUTHORS Double-strand nucleic acid probes and the use thereof
TITLE Patent: WO 0131057-A 3 03-MAY-2001
JOURNAL Aventis Research & Technologies GmbH & Co KG (DE)
FEATURES Location/Qualifiers
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Qy 1 ATTCGATCGGGCGGGCGGAGC 22
Db 49 ATTCGATCGGGCGGGCGGAGC 28

RESULT 12
AX377573
LOCUS AX377573 49 bp DNA linear PAT 18-MAR-2002
DEFINITION Sequence 50 from Patent WO0212553.
ACCESSION AX377573
VERSION AX377573.1 GI:19573759
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1 Kappel, A., Polakowski, T., Pignot, M., Windhab, N., Behrens, H., and Muth, J.
AUTHORS Method for detecting mutations in nucleotide sequences
TITLE Patent: WO 0212553-A 50 14-FEB-2002;
JOURNAL Nanogen Recognomics GmbH (DE)
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Hairpin-Oligonucleotid"
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BASE COUNT 6 a 14 c 16 g 13 t
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Query Match 100.0%; Score 22; DB 6; Length 49;
Best Local Similarity 100.0%; Pred. No. 45;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAGC 22
Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 13
AX104633
LOCUS AX104633 21 bp DNA linear PAT 30-APR-2001
DEFINITION Sequence 825 from Patent WO0122972.
ACCESSION AX104633
VERSION AX104633.1 GI:13920830
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1 Krieg, A.M., Schetter, C. and Vollmer, J.C.
AUTHORS Immunostimulatory nucleic acids
TITLE Patent: WO 0122972-A 825 05-APR-2001;
JOURNAL UNIVERSITY OF IOWA RESEARCH FOUNDATION (US); Coley Pharmaceutical GmbH (DE)
FEATURES Location/Qualifiers
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BASE COUNT 3 a 4 c 11 g 3 t
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Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 ATTCGATCGGGCGGGCGGAGC 21

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RESULT 14
AX104634/c
LOCUS AX104634 21 bp DNA linear PAT 30-APR-2001
DEFINITION Sequence 826 from Patent WO0122972.
ACCESSION AX104634
VERSION AX104634.1 GI:13920831
KEYWORDS
ORGANISM synthetic construct
SOURCE synthetic construct
REFERENCE 1
AUTHORS Krieg,A.M.; Schetter,C. and Vollmer,J.C.
TITLE Immunostimulatory nucleic acids
JOURNAL Patent: WO 0122972-A 826 05-APR-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US) ; Coley Pharmaceutical
GmbH (DE)
FEATURES
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3 a 11 c 4 g 3 t
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Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 21 ATTCGATCGGGCGGGCGGAG 1

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AX355087
LOCUS AX355087 21 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 115 from Patent WO0197843.
ACCESSION AX355087
VERSION AX355087.1 GI:18619754
KEYWORDS
ORGANISM synthetic construct
SOURCE synthetic construct
REFERENCE 1
AUTHORS Weiner,G. and Hartmann,G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
JOURNAL cancer
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)
FEATURES
Source
Location/Qualifiers
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/organism="synthetic construct"
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BASE COUNT
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Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 16
AX355201/c
LOCUS AX355201 21 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 229 from Patent WO0197843.
ACCESSION AX355201
VERSION AX355201.1 GI:18619868

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KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Weiner,G. and Hartmann,G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
JOURNAL cancer
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)
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Source
Location/Qualifiers
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/mol_type="genomic DNA"
/db_xref="taxon:32630"
3 a 11 c 4 g 3 t
BASE COUNT
3 a 11 c 4 g 3 t
ORIGIN
Query Match 95.5%; Score 21; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 17
AX547686
LOCUS AX547686 21 bp DNA linear PAT 26-NOV-2002
DEFINITION Sequence 825 from Patent WO02053141.
ACCESSION AX547686
VERSION AX547686.1 GI:25812830
KEYWORDS
ORGANISM synthetic construct
SOURCE synthetic construct
REFERENCE 1
AUTHORS Bratzler,R.L.
TITLE Inhibition of angiogenesis by nucleic acids
JOURNAL Patent: WO 02053141-A 825 11-JUL-2002;
Coley Pharmaceutical Group, Inc. (US)
FEATURES
Source
Location/Qualifiers
1..21
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
3 a 4 c 11 g 3 t
BASE COUNT
3 a 4 c 11 g 3 t
ORIGIN
Query Match 95.5%; Score 21; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
Db 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 18
AX547687/c
LOCUS AX547687 21 bp DNA linear PAT 26-NOV-2002
DEFINITION Sequence 826 from Patent WO02053141.
ACCESSION AX547687
VERSION AX547687.1 GI:25812831
KEYWORDS
ORGANISM synthetic construct
SOURCE synthetic construct
REFERENCE 1
AUTHORS Bratzler,R.L.

```


TITLE Inhibition of angiogenesis by nucleic acids
JOURNAL Patent: WO 0203141-A 826 11-JUL-2002;
Coley Pharmaceutical Group, Inc. (US)

FEATURES
source
1. .21
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/note="Synthetic Sequence"

BASE COUNT 3 a 11 c 4 g 3 t

ORIGIN

Query Match 95.5%; Score 21; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21

Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 19

A89791
LOCUS A89791 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 13 from Patent WO9832462.
ACCESSION A89791
VERSION A89791.1 GI:6738305

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

Patent: WO 9832462-A 13 30-JUL-1998;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)

FEATURES

source

1. .20
/organism="unidentified"

/mol_type="genomic DNA"

/db_xref="taxon:32644"

BASE COUNT 2 a 5 c 11 g 2 t

ORIGIN

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAG 22

Db 1 TCGATCGGGCGGGCGGAG 20

RESULT 20

A90878
LOCUS A90878 20 bp DNA linear PAT 22-JAN-2000
DEFINITION Sequence 13 from Patent EP0855184.
ACCESSION A90878
VERSION A90878.1 GI:6739281

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

Patent: EP 0855184-A 13 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)

FEATURES

source

1. .20
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/mol_type="genomic DNA"

/db_xref="taxon:32630"

/note="Synthetic oligonucleotide"

BASE COUNT 2 a 5 c 11 g 2 t

ORIGIN

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/db_xref="taxon:32644"

BASE COUNT 2 a 5 c 11 g 2 t

ORIGIN

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAG 22

Db 1 TCGATCGGGCGGGCGGAG 20

RESULT 21

AX023404
LOCUS AX023404 20 bp DNA linear PAT 15-SEP-2000
DEFINITION Sequence 19 from Patent WO0014217.
ACCESSION AX023404
VERSION AX023404.1 GI:10183804

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

Lipford, G.B., Heeg, K. and Wagner, H.
G-motif oligonucleotides and uses thereof
Patent: WO 0014217-A 19 16-MAR-2000;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE); WAGNER HERMANN (DE);
CPG IMMUNOPHARMACEUTICALS GMBH (DE)

FEATURES

source

1. .20
/organism="synthetic construct"

/mol_type="genomic DNA"

/db_xref="taxon:32630"

/note="synthetic, no natural origin"

BASE COUNT 2 a 5 c 11 g 2 t

ORIGIN

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAG 22

Db 1 TCGATCGGGCGGGCGGAG 20

RESULT 22

AX455627
LOCUS AX455627 20 bp DNA linear PAT 06-JUL-2002
DEFINITION Sequence 104 from Patent WO0222809.
ACCESSION AX455627
VERSION AX455627.1 GI:21714695

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

Bauer, S., Lipford, G. and Wagner, H.
Process for high throughput screening of cpg-based
immuno-agonist/antagonist
Patent: WO 0222809-A 104 21-MAR-2002;
Coley Pharmaceutical GmbH (DE)

FEATURES

source

1. .20
/organism="synthetic construct"

/mol_type="genomic DNA"

/db_xref="taxon:32630"

/note="Synthetic oligonucleotide"

BASE COUNT 2 a 5 c 11 g 2 t

ORIGIN

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 1 TCGATCGGGCGGGCGGAGC 20

RESULT 23
LOCUS BD056782 20 bp DNA linear PAT 27-AUG-2002
DEFINITION Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination.
ACCESSION BD056782
VERSION JP 2001508780-A/12
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1 (bases 1 to 20)
AUTHORS Wagner, H., Lipford, G.B. and Heeg, K.
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination
JOURNAL Patent: JP 2001508780-A 12 03-JUL-2001;
COMMENT HERMANN WAGNER, GRAYSON B LIPFORD, KLAUS HEEG
PD 03-JUL-2001
PF 23-JAN-1998 JP 1998531592
PR 23-JAN-1997 EP 97101019.4
PI HERMANN WAGNER, GRAYSON B LIPFORD, KLAUS HEEG
PC A61K39/39, A61K31/7088, A61K39/00, A61P37/04, C12N15/09, C12N15/00
CC Strandedness: Single;
CC Topology: Linear;
FH Key Location/Qualifiers.
FEATURES
source 1..20
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
BASE COUNT 2 a 5 c 11 g 2 t
ORIGIN

Query Match 90.9%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 1 TCGATCGGGCGGGCGGAGC 20

RESULT 24
LOCUS BD056782 46 bp DNA linear PAT 03-APR-1998
DEFINITION Sequence 12 from patent US 5683985.
ACCESSION I72381
VERSION I72381.1 GI:3008520
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 46)
AUTHORS Chu, B.Chen, Fei, and Orgel, L.
TITLE Oligonucleotide decoys and methods relating thereto
JOURNAL Patent: US 5683985-A 12 04-NOV-1997;
FEATURES
source 1..46
/organism="unknown"
BASE COUNT 3 a 21 c 15 g 7 t
ORIGIN

Query Match 86.4%; Score 19; DB 6; Length 46;
Best Local Similarity 100.0%; Pred. No. 8.5e+02;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Best Local Similarity 100.0%; Pred. No. 8.5e+02;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 21
DB 28 TCGATCGGGCGGGCGGAGC 46

RESULT 25
LOCUS I72382 46 bp DNA linear PAT 03-APR-1998
DEFINITION Sequence 13 from patent US 5683985.
ACCESSION I72382
VERSION I72382.1 GI:3008521
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 46)
AUTHORS Chu, B.Chen, Fei, and Orgel, L.
TITLE Oligonucleotide decoys and methods relating thereto
JOURNAL Patent: US 5683985-A 13 04-NOV-1997;
FEATURES
source 1..46
/organism="unknown"
BASE COUNT 3 a 21 c 15 g 7 t
ORIGIN

Query Match 86.4%; Score 19; DB 6; Length 46;
Best Local Similarity 100.0%; Pred. No. 8.5e+02;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 21
DB 28 TCGATCGGGCGGGCGGAGC 46

RESULT 26
LOCUS AX377573 49 bp DNA linear PAT 18-MAR-2002
DEFINITION Sequence 50 from Patent WO0212553.
ACCESSION AX377573
VERSION AX377573.1 GI:19573759
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1
AUTHORS Kappel, A., Polakowski, T., Pignot, M., Windhab, N., Behrens, H. and Much, J.
TITLE Method for detecting mutations in nucleotide sequences
JOURNAL Patent: WO 0212553-A 50 14-FEB-2002;
Nanogen Recognomics GmbH (DE)
FEATURES
source 1..49
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/note="Beschreibung der künftlichen Sequenz: Hairpin-Oligonucleotid"
BASE COUNT 6 a 14 c 16 g 13 t
ORIGIN

Query Match 85.5%; Score 18.8; DB 6; Length 49;
Best Local Similarity 90.9%; Pred. No. 1e+03;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 49 ATTCGATCGGGCGGGCGGAGC 28

RESULT 27

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E07877
LOCUS       E07877                22 bp    DNA    linear    PAT 29-SEP-1997
DEFINITION   Synthetic nucleotides for DNA binding protein analysis.
ACCESSION    E07877
VERSION      E07877.1 GI:2176010
KEYWORDS     JP 1994201692-A/2.
SOURCE       unidentified
ORGANISM     unidentified
REFERENCE    1 (bases 1 to 22)
AUTHORS      Uno,I., Ichihara,T., Igarashi,K., Miki,N. and Ous.S.
TITLE        PROBE FOR DETECTING OR MEASURING DNA-BINDING PROTEIN AND METHOD FOR
JOURNAL      DETECTING OR MEASURING DNA-BINDING PROTEIN USING IT
PATENT       Patent: JP 1994201692-A 2 22-JUL-1994;
NIPPON STEEL CORP, NIPPON STEEL CHEM CO LTD
OS           None
OC           Artificial sequences.
PN           JP 1994201692-A/2
PD           22-JUL-1994
PF           22-SEP-1993 JP 1993257482
PR           22-SEP-1992 JP 92P 278126
PI           UNO ISAO, ISOHARA TOSHIO, IGARASHI KATSUHIKO, MIKI NAOMASA, PI
PC           GOIN33/53//C12Q1/69;
CC           strandedness: Single;
CG           topology: Linear;
CH           Location/Qualifiers
FH           key
FT           source
PT           1..22
              Location/Qualifiers
FEATURES     source
             1..22
             /organism="Artificial sequences".
BASE COUNT   3 a      11 g      3 t
ORIGIN
1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
TCGATCGGGGGCGGGCGGAGC 22
|||||
TCGATCGGGGGCGGGCGGATC 22

Query Match      83.6%; Score 18.4; DB 6; Length 22;
Best Local Similarity 95.0%; Pred. No. 1.6e+03;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY
3 TCGATCGGGGGCGGGCGGAGC 22
|||||
3 TCGATCGGGGGCGGGCGGATC 22

Db

RESULT 28
LOCUS       172381/c
DEFINITION   Sequence 12 from patent US 5683985.
ACCESSION    172381
VERSION      172381.1 GI:3008520
KEYWORDS
SOURCE       Unknown.
ORGANISM     Unclassified.
REFERENCE    1 (bases 1 to 46)
AUTHORS      Chu,B.Chen.Fei. and Orgel,L.
TITLE        Oligonucleotide decoys and methods relating thereto
JOURNAL      Patent: US 5683985-A 12 04-NOV-1997;
FEATURES     Location/Qualifiers
             source
             1..46
             /organism="unknown"
BASE COUNT   3 a      21 c      15 g      7 t
ORIGIN
1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
GATCGGGGGCGGGCGGAGC 22
|||||
GATCGGGGGCGGGCGGAGC 22

Query Match      81.8%; Score 18; DB 6; Length 46;
Best Local Similarity 100.0%; Pred. No. 2.3e+03;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
5 GATCGGGGGCGGGCGGAGC 22
|||||
GATCGGGGGCGGGCGGAGC 22

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```

Db
24 GATCGGGGGCGGGCGGAGC 7

RESULT 29
LOCUS       172382/c
DEFINITION   Sequence 13 from patent US 5683985.
ACCESSION    172382
VERSION      172382.1 GI:3008521
KEYWORDS
SOURCE       Unknown.
ORGANISM     Unclassified.
REFERENCE    1 (bases 1 to 46)
AUTHORS      Chu,B.Chen.Fei. and Orgel,L.
TITLE        Oligonucleotide decoys and methods relating thereto
JOURNAL      Patent: US 5683985-A 13 04-NOV-1997;
FEATURES     Location/Qualifiers
             source
             1..46
             /organism="unknown"
BASE COUNT   3 a      21 c      15 g      7 t
ORIGIN
1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
GATCGGGGGCGGGCGGAGC 22
|||||
GATCGGGGGCGGGCGGAGC 7

Query Match      81.8%; Score 18; DB 6; Length 46;
Best Local Similarity 100.0%; Pred. No. 2.3e+03;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
5 GATCGGGGGCGGGCGGAGC 22
|||||
GATCGGGGGCGGGCGGAGC 7

Db

RESULT 30
LOCUS       AC144487/c
DEFINITION   Gasterosteus aculeatus clone CH213-16009, complete sequence.
ACCESSION    AC144487
VERSION      AC144487.1 GI:30089725
KEYWORDS     HTG.
SOURCE       Gasterosteus aculeatus (three spined stickleback)
ORGANISM     Gasterosteus aculeatus (three spined stickleback)
REFERENCE    1 (bases 1 to 181162)
AUTHORS      Kingsley,D., Peichel,C., Colosimo,P., Shapiro,M., Marks,M.,
Blackman,B., Grimwood,J., Dickson,M., Schmutz,J. and Myers,R.M.
TITLE        Genomic studies of vertebrate diversity
JOURNAL      Unpublished
AUTHORS      2 (bases 1 to 181162)
REFERENCE    Kingsley,D., Peichel,C., Colosimo,P., Shapiro,M., Marks,M.,
Blackman,B., Grimwood,J., Dickson,M., Schmutz,J. and Myers,R.M.
TITLE        Direct Submission
JOURNAL      Submitted (24-APR-2003) Stanford Human Genome Center, 975
              California Avenue, Palo Alto, CA 94304, USA
REFERENCE    3 (bases 1 to 181162)
AUTHORS      Kingsley,D., Peichel,C., Colosimo,P., Shapiro,M., Marks,M.,
Blackman,B., Grimwood,J., Dickson,M., Schmutz,J. and Myers,R.M.
TITLE        Direct Submission
JOURNAL      Submitted (17-MAY-2003) Stanford Human Genome Center, 975
              California Avenue, Palo Alto, CA 94304, USA
COMMENT      The sequence of the clone was established as a mapping and
              sequencing collaboration at the Stanford Genome Evolution Center,
              funded by the NIH Centers of Excellence in Genomic Science (CEGS),
              the BAC library CHORI213 built by Pieter deon in collaboration
              with the Stanford Genome Evolution Center
              (http://www.chori.org/bacpac/). Further details on the source fish
              and construction of the library can be found at
              http://cegs.stanford.edu/stickleback/BAC_library_link.jsp.
              Quality: Phrap Quality >=40 100% of Sequence;
              Estimated Total Number of Errors is 0.

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FEATURES
  source      Location/Qualifiers
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              /organism="Gasterosteus aculeatus"
              /mol_type="genomic DNA"
              /db_xref="taxon:69293"
              /clone="CH213-16009"
BASE COUNT   52530 a 39127 c 37839 g 51666 t
ORIGIN
Query Match      80.9%; Score 17.8; DB 5; Length 181162;
Best Local Similarity 90.5%; Pred. No. 1.6e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGCGAG 21
    |||||
Db 150119 ATTCGATCGGGCGGGCGGCGG 150099

RESULT 31
AC127795      274015 bp      DNA      linear      HTG 20-NOV-2002
LOCUS         Rattus norvegicus clone CH230-101A9, *** SEQUENCING IN PROGRESS
DEFINITION    ***. 3 unordered pieces.
ACCESSION     AC127795.3 GI:25139837
VERSION       HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.
KEYWORDS      Rattus norvegicus (Norway rat)
SOURCE        Rattus norvegicus
ORGANISM      Eukaryota; Metazoa; Chordata; Vertebrata; Eureleostomi;
              Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
              Rattus.
REFERENCE
  1 (Bases 1 to 274015)
  Muzny, D., Marie, E., Metzker, M., Lee, S., Abramson, S., Adams, C., Alder, J.,
  Allen, C., Allen, H., Alsbrooks, S., Amin, A., Angiano, D.,
  Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
  Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,
  Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
  Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,
  Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,
  Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,
  Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
  Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,
  Delgado, O., Denison, S., Deramo, C., Ding, Y., Dinh, H., Diya, K.,
  Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,
  Egan, A., Escotto, M., Eugene, C., Evans, C., Falls, T., Fan, G.,
  Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,
  Fraser, C., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M.,
  Gebregorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W.,
  Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K.,
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  Hernandez, R., Hines, S., Hladun, S., Hodgson, A., Hogue, M.,
  Hollins, B., Howells, S., Huylk, S., Hume, J., Idlebird, D., Jackson, A.,
  Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,
  Karpachy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C.,
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  Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,
  Lorenshewa, L., Loulseg, H., Lozada, R., Lu, X., Ma, J.,
  Maheshwari, M., Mahindratne, M., Mahmoud, M., Mallory, K., Mangum, A.,
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  Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H.,
  Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C., Smajs, D.,
  Sneed, A., Sodergren, E., Song, X., Sorelle, R., Sosa, J.,
  Steilmie, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C.,

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Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K.,
Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J.,
Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,
Williams, G., Willson, R., Wleczyk, R., Wooden, H., Worley, K.,
Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,
Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von
Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O.,
Weinstock, G. and Gibbs, R.A.
Direct Submission
Unpublished
2 (bases 1 to 274015)
Worley, K.C.
Direct Submission
Submitted (19-JUL-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 274015)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (20-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Nov 20, 2002 this sequence version replaced gi:23269472.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center of Medicine
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GZMW
Center clone name: CH230-101A9
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 219062 bases at least Q40
Consensus quality: 22791 bases at least Q30
Consensus quality: 225601 bases at least Q20
Estimated insert size: 225142; sum-of-contigs estimation
Quality coverage: 5x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
consists of 3 contigs. The true order of the pieces
is not known and their order in this sequence record is
arbitrary. Gaps between the contigs are represented as
runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
as soon as it is available and the accession number will
be preserved.
-----
1 9143: contig of 9143 bp in length
9144 9243: gap of unknown length
9244 272830: contig of 263587 bp in length
272831 272930: gap of unknown length
272931 274015: contig of 1085 bp in length.
Location/Qualifiers
1. 274015
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-101A9"
1. 1184
misc_feature

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misc_feature /note="wgs contig"
 9244. .11119
 /note="wgs_contig"
 BASE COUNT 61069 a 49181 c 50666 g 66519 t 46580 others
 ORIGIN

Query Match 80.9%; Score 17.8; DB 2; Length 274015;
 Best Local Similarity 90.5%; Pred.No.1.5e+03;
 Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 ATTCCATCGGCGGGCGGAG 21
 |||||

Db 228660 ATTCCATCGGCGGGCGGAG 228680

RESULT 32

AC112802
 LOCUS
 DEFINITION Rattus norvegicus clone CH230-35H17, *** SEQUENCING IN PROGRESS
 AC112802
 AC112802.3 GI:23101275
 HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ENRICHED.
 VERSION
 KEYWORDS Rattus norvegicus (Norway rat)
 SOURCE Rattus norvegicus
 ORGANISM
 Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
 Rattus.
 1 (bases 1 to 301660)
 REFERENCE
 AUTHORS
 Muzny,D,Marie, Metzker,M, Lee, A, Adams, C., Alder, J.,
 Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
 Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
 Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,
 Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
 Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,
 Cardenas, V., Carter, K., Cavazos, I., Cesar, H., Center, A., Chu, J.,
 Chacko, J., Chavez, D., Chen, R., Chen, Y., Chen, Z., Chu, J.,
 Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
 Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,
 Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,
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 Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G.,
 Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,
 Frazer, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, I., Garza, M.,
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 Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K.,
 Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J.,
 Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hognes, M.,
 Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A.,
 Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,
 Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C.,
 Kowis, C., Kratt, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J.,
 Liu, J., Liu, W., Liu, X., London, P., Longacre, S., Lopez, J.,
 Lorenshewa, L., Loulseghe, H., Lozado, R.J., Lu, X., Ma, J.,
 Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A.,
 Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E.,
 Mawhiney, S., McLeod, M.P., McNeill, T.Z., Meenen, E.,
 Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,
 Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L.,
 Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S.,
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 Pasternak, S., Paul, H., Perez, A., Perez, L., Pfankoch, C.,
 Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.,
 Puato, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R.,
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 Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J.,
 Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H.,
 Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajls, D.,
 Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J.,
 Steinle, M., Strong, R., Sutton, A., Svatek, A., Tabot, P., Taylor, C.,
 Taylor, R., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K.,
 Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J.,
 Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,

Williams, G., Willson, R., Wlarczyk, R., Wooden, H., Worley, K.,
 Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,
 Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von
 Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O.,
 Weinstein, G. and Gibbs, R.A.
 Direct Submission
 Unpublished
 2 (bases 1 to 301660)
 Worley, K.C.
 Direct Submission
 Submitted (25-FEB-2002) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 3 (bases 1 to 301660)
 Rat Genome Sequencing Consortium.
 Direct Submission
 Submitted (22-SEP-2002) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 On Sep 18, 2002 this sequence version replaced gi:21737447.
 The sequence in this assembly is a combination of BAC based reads
 and whole genome shotgun sequencing reads assembled using Atlas
 (http://www.hgsc.bcm.tmc.edu/projects/rat/). As a result, the
 sequence may extend beyond the ends of the clone and there may be
 contigs that consist entirely of whole genome shotgun sequence
 reads. Both end sequences and whole genome shotgun sequence only
 contigs will be indicated in the feature table.
 ----- Genome Center
 Center: Baylor College of Medicine
 Center code: BCM
 Web site: http://www.hgsc.bcm.tmc.edu/
 Contact: hgsc-help@bcm.tmc.edu
 ----- Project Information
 Center project name: GPD
 Center clone name: CH230-35H17
 ----- Summary Statistics
 Assembly program: Phrap; version 0.990329
 Consensus quality: 228218 bases at least Q40
 Consensus quality: 231561 bases at least Q30
 Consensus quality: 233804 bases at least Q20
 Estimated insert size: 253557; sum-of-contigs estimation
 Quality coverage: 4x in Q20 bases; sum-of-contigs estimation

 * NOTE: Estimated insert size may differ from sequence length
 * (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)
 * NOTE: This sequence may represent more than one clone.
 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 2 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.
 * 1 197738: contig of 197738 bp in length
 * 197739 197838: gap of unknown length
 * 197839 301660: Contig of 103822 bp in length.

FEATURES

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 /db_xref="taxon:10116"
 /clone="CH230-35H17"
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 1..1034
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 5898..6267
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 /note="T7
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 end_sequence:BH276513"
 6224..6827
 /note="clone_boundary
 misc_feature
 6224..6827
 /note="clone_boundary

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site:ECORI
end_sequence:BH276515"
misc_feature
197839..199723
/notes="wgs_end_extension
clone_end:sp6"
BASE COUNT 67072 a 49216 c 49408 g 69772 t 66192 others
ORIGIN
Query Match 80.9%; Score 17.8; DB 2; Length 301660;
Best Local Similarity 90.5%; Pred. No. 1.5e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
Db 15144 ATTCGATCGGGCGGGCGGAG 15164

RESULT 33
BC016449
LOCUS
DEFINITION Mus musculus ring finger protein 5, mRNA (cdna clone MGC:19320
IMAGE:4191746), complete cds.
ACCESSION BC016449
VERSION BC016449.1 GI:16741215
SOURCE MGC.
ORGANISM Mus musculus (house mouse)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 1119)
Straussberg,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G.,
Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
Altschul,S.F., Zuber,B., Buetow,K.H., Schaefer,C.F., Bhat,N.K.,
Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Hsieh,F.,
Diatchenko,L., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
Stapleton,M., Soares,M.B., Bonaldo,M.F., Casavant,T.L.,
Schetz,T.E., Brownstein,M.J., Usdin,T.B., Toshiyuki,S.,
Carninci,P., Frange,C., Raja,S.S., Loquellano,N.A., Peters,G.J.,
Abramson,R.D., Mallek,S.J., Bosak,S.A., McEwan,P.J.,
McKernan,K.J., Mullik,J.A., Gunaratne,P.H., Richards,S.,
Worley,K.C., Hale,S., Garcia,A.M., Gay,L.J., Hulyk,S.W.,
Villalon,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A.,
Fahey,J., Helton,E., Kerteman,M., Madan,A., Rodrigues,S.,
Sanchez,A., Whitting,M., Madan,A., Young,A.C., Shevchenko,Y.,
Bouffard,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D.,
Dickson,M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M.,
Butterfield,Y.S., Krzywinski,M.I., Skalska,U., Smalhus,D.E.,
Schnerch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences
Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
22388257
12477932
2 (bases 1 to 1119)
Straussberg,R.
Direct Submission
Submitted (31-OCT-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: cgaaps@mail.nih.gov
Tissue Procurement: Jeffrey E. Green, M.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Baylor College of Medicine Human Genome
Sequencing Center
Center code: BCM-HGSC
Web site: http://www.hgsc.bcm.tmc.edu/cdna/
Contact: amg@bcm.tmc.edu
Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Louised, H.,

```

Kowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Nanavati, A.N., Gibbs, R.A.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
 Series: IRAK Plate: 24 Row: i Column: 17
 This clone was selected for full length sequencing because it passed the following selection criteria: Hexamer frequency ORF analysis, GenomeScan gene prediction, similarity but not identity to protein.

Location/Qualifiers
 1..1119
 /organism="Mus musculus"
 /mol_type="mRNA"
 /strain="FVB/N"
 /db_xref="taxon:10090"
 /clone="MGC:19320 IMAGE:4191746"
 /tissue_type="Salivary gland, 10 week old female mouse"
 /clone_lib="NCI CGAP_SG2"
 /lab_host="DH10B"
 /note="Vector: pCMV-SPORT6"
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 /gene="Rnf5"
 /note="synonyms: 2410131005RIK, NG2"
 /db_xref="LocusID:54197"
 /db_xref="MGI:1860076"
 89..631
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 /db_xref="GI:16741216"
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 LYWPCLHOLWTRPDRQPCVCKAGISREKVLYVGRGSKQPDRLKTPPEQQR
 PAGESGGFQPGDAGHEHESFGVGAFPGFTTVAHEPFRGAGVLDGQGHPRAS
 WQSLSEFLAIFFFWLLSI"

BASE COUNT 244 a 292 c 296 g 287 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 10; Length 1119;
 Best Local Similarity 94.7%; Pred. No. 3.3e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAG 22

Db 136 CGAGCGGGCGGGCGGCGAG 154

RESULT 34

LOCUS

DEFINITION HSA243936 Homo sapiens mRNA for Gl6 protein (Gl6 gene located in the class III region of the major histocompatibility complex).

ACCESSION AJ243936

VERSION AJ243936.1 GI:5578772

KEYWORDS Gl6 gene; Gl6 protein.

SOURCE Homo sapiens

ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 Kendall,E., Sargent,C.A. and Campbell,R.D.

AUTHORS Human major histocompatibility complex contains a new cluster of genes between the HLA-D and complement C4 loci

JOURNAL Nucleic Acids Res. 18 (24), 7251-7257 (1990)

MEDLINE 91081311

PUBMED 2259622

REFERENCE 2 Khanna,A.

AUTHORS Thesis (1993) University of Oxford, Department of Biochemistry. MRC Immunochimistry Unit

JOURNAL Khanna,A. and Campbell,R.D.

REFERENCE 3

TITLE Characterisation of a novel gene, G16, in the class III region of the human Major Histocompatibility Complex

JOURNAL Unpublished

REFERENCE 4 (bases 1 to 1123)

AUTHORS Aguado B.

JOURNAL Direct Submission

TITLE Submitted (20-JUL-1999) Aguado B., HGMP Resource Centre, MEC, Genome Campus, Hinxton, Cambridge, CB10 1SB, UNITED KINGDOM

FEATURES

source

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/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/chromosomes="6"

/map="6p21.3"

/cell_line="U937"

/notes="class III region of the major histocompatibility complex"

gene

1..1123

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/function="unknown"

/notes="contains a Ring Finger"

/codon_start=1

/product="protein G16"

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/db_xref="GI:5578773"

/translations="MAAEEDEGGPGPNRGGASATFECNICLETAREAVSVCGH LYCWPCLEHWRLETRPQPCVCVKGISREKVVPLIGRSGRQKQDFRLKTPRPGQR PAPESGGFQPGDAGGFFHFGVAGFPFGFTTFVNAHEPPRRGAGVLDGGHPPAS WQSLFLFLAIFPFWLLSI"

110..1105

/gene="G16"

/note="putative"

978..982

/gene="G16"

/notes="mRNA instability motif (AUUUA)"

1088..1093

/gene="G16"

1094..1099

/gene="G16"

/notes="putative"

BASE COUNT 240 a 292 c 299 g 292 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 9; Length 1123;

Best Local Similarity 94.7%; Pred. No. 3.3e+03;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGGGGGCGGCGGAGC 22

Db 148 CGAGCGGGGGCGGCGGAGC 166

RESULT 35

AF335479

LOCUS AF335479 7125 bp DNA linear BCT 02-FEB-2002

DEFINITION Agrobacterium sp. IP I-671 putative transposase gene, partial cds; hydantoin utilization gene cluster, complete sequence; and putative resolvase gene, partial cds.

ACCESSION AF335479

VERSION AF335479.1 GI:18478559

KEYWORDS

SOURCE

ORGANISM

Agrobacterium sp. IP I-671

Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales; Rhizobiaceae; Rhizobium/Agrobacterium group; Agrobacterium.

1 (bases 1 to 7125)

REFERENCE

AUTHORS Hils, M. and Altenbuchner, J.

TITLE Hydantoin utilization genes of Agrobacterium sp. IP I-671

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 7125)

AUTHORS Hils M. and Altenbuchner, J.

TITLE Direct Submission

JOURNAL Submitted (11-JAN-2001) Institute of Industrial Genetics, University of Stuttgart, Almandring 31, Stuttgart 70569, Germany

FEATURES

source

1..7125

/organism="Agrobacterium sp. IP I-671"

/mol_type="genomic DNA"

/strain="IP I-671"

/db_xref="taxon:173261"

/complement("1..445)

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/db_xref="GI:18478560"

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complement (576..1286)

/gene="hyuN"

/complement (576..1286)

/gene="hyuN"

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/product="putative NADPH dependent flavin oxidoreductase"

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/db_xref="GI:18478561"

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complement (1311..2684)

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/note="hyuH"

/codon_start=1

/transl_table=11

/product="D-hydantoinase"

/protein_id="AAL73199.1"

/db_xref="GI:18478562"

/translations="MDIILKNGITVADGISRADIGIKGKIIVGIGALGAERTIDA SGRYVPGVDVHTVETSFNTCSADTFATVAAACGGTTIIVDFCQDGRSLTD AVAKWDGAGKGAIDYGYHII VLDPTDSVIELEVPGLTISFKVFMAYRGNMID DVTLLKTDKAARTGSLVMHAENGDAADYLRNKVAGETAPIYHALSPPRIEAEA TARALAEIVDAPYIVHTVTCESLDEVMRSKARGVHALAETCTHYLYLKEDLERP GFEGAKYVTPPARAKDHEILWNALRNGAFETVSSDHCWMLFRGHDKGRNDRAIP NGAGVEERLMVYQGVNEGRLSITQFVELVATRPAAVFGMFPEKGTIAGSDADI VWPRAEMVIEQSAMHNDYSTVEGRKVKGVPTVLLRGRVIVEDSGSYVCAPTDGGFL KRRRYKQ"

3154..4068

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3154..4068

/gene="hyuC"

/note="hyuC"

/codon_start=1

/transl_table=11

/product="D-carbamoylase"

/protein_id="AAL73200.1"

/db_xref="GI:18478563"

/translations="WTRQMLAVGQGGPIARAETREQVIARLLDMLANAASRGVNFIV FPELAVTFFPRWLTDEAEDLSFVETEMPGLTRPLFEKAAELGIGNFYGAELVVE CGVRRFNTSILVDRSGKIGIKYKVKHLPQHKVEAYRPPQHLKRYFEPDGMGFPVY DVDAAKMGWFCNDRRMPAEWRVNLKGAELICGGYNTPTNPAVPOHDLTSPHLL SQAGSYQNGASAAAGKVGMEEDMLLGHSCIVAPTGEIVALTITLEDDEVITAMIDL DRCELRREHIFNKAHQPOHYGLIAEL"

4085..5338

/gene="hyuB"

4085..5338

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CDS

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/notes="hyuD"
/codon_start=1
/transl_table=11
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GQISPLSPFAPFGLVKKAVHAFAYEPFLIVGRVDPDAMARFLWRWRASSVDQIV
TAGKAVTIAEYSDRCFALRREIPIDYAGRGRLVAFSPQSDGVDGAVDLAVDEL
RVPYRLREAEVAFENFIAPTSVGIVGQLETSDEGCFRETQALACRAENGVRVY
YGAASALLMENGRIKAVTAEADVEMDAVVVALGVWSNALLPLGIDLPYIPVKGY
LTVKADPDTFGMATISDTKYGVTNLGDIRVGTGAELAGFVDSQPEKRVAGLYOT
LKSLEPKVPEAAIADAERNSGLRPVTPDGPPLIGRSRIPNLVINSHGHTLGTWNSCGS
GRLADVIGDERPAVLSRPAFTAL"
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/notes="hyuA"
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/product="hydantoin-racemase"
/protein_id="AAL73202.1"
/db_xref="GI:18478565"
/translation="MKIKVINPTNTWTMDKLAGAARAAAAPGTEIVAVSPDMGPVSI
EGYIDVFAVGVDEVRKGELEGCDGVYIACGDPGLNAAREVARGVIGIAEAMH
VLEEGSGFSIISMLGRSGVLEHLVHVSXGMAHKCRVMTDLFVLEFEESGDARRI
VVEECRAIEQDHASVILGCGSMDLMAVYSGEIGAPALDGVSGVXLVEALVGWGL
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6875..>7125
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/transl_table=11
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/db_xref="GI:18478566"
/translation="MLIGMYRVSQDERQSVLQRDALIAAGVDQRLHQDRASGARD
DRPGLKACIAECEDGLVWKLDRGLSLHLRIVEDP"
BASE COUNT 1403 a 2103 c 2146 g 1473 t
ORIGIN
Query Match 79.1%; Score 17.4; DB 1; Length 7125;
Best Local Similarity 94.7%; Pred. No. 2.9e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
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Db 902 CGATCGGGCGGGCGGAGC 920

RESULT 36
AX346590
LOCUS AX346590 9646 bp DNA linear PAT 01-FEB-2002
DEFINITION Sequence 1661 from Patent WO0200928.
ACCESSION AX346590
VERSION AX346590.1 GI:18494476
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.
TITLE Diagnosis of diseases associated with the immune system
JOURNAL Patent: WO 0200928-A 1661 03-JAN-2002;
Epigenomics AG (DE)
FEATURES
source
Location/Qualifiers
1..9646
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/notes="chemically treated genomic DNA (Homo sapiens)"
BASE COUNT 2765 a 150 c 2136 g 4595 t
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ORIGIN

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Query Match 79.1%; Score 17.4; DB 6; Length 9646;
Best Local Similarity 94.7%; Pred. No. 2.8e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 21
|||||
Db 5203 TCGATCGGGCGGGCGGAGC 5221
```

RESULT 37

```
HSLUCAL/c
LOCUS HSLUCAL 30377 bp DNA linear PRI 04-MAR-2003
DEFINITION Human DNA sequence from clone XXcos-1 on chromosome 3, complete
sequence.
ACCESSION Z74618
VERSION Z74618.1 GI:1405891
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
```

REFERENCE

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AUTHORS


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QY      4 CGATCGGGCGGGCGGAGC 22
Db      29479 CGAGCGGGCGGGCGGAGC 29461

RESULT 38
RN520N23/c
LOCUS   106278 bp DNA linear HTG 04-JUN-2002
DEFINITION Rattus norvegicus clone RPCI-31-520N23 strain Brown Norway, ***
SEQUENCING IN PROGRESS ***, 138 unordered pieces.
ACCESSION AL732653.1 GI:21326744
VERSION   HTG; HTGS PHASE1; HTGS DRAFT.
KEYWORDS  Rattus norvegicus (Norway rat)
SOURCE   Rattus norvegicus
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE 1
AUTHORS  Sudbrak,R., Borzym,K., Mueller,I., Klages,S., Kosiura,A.,
Walter,L., Guenther,E., Hurt,P., Lehrach,H., Himmelbauer,H. and
Reinhardt,R.
JOURNAL  Unpublished
REFERENCE 2 (bases 1 to 106278)
AUTHORS  MOLGENR.
JOURNAL  Direct Submission
TITLE    Submitted (04-JUN-2002) MPIMG, Abt. Lehrach, Max Planck Institut
Fuer Molekulare Genetik, Ihnestrasse 73, Berlin, 14195 Germany
COMMENT  1. 753
contig 01 854..1659
contig 02 1760..2272
contig 03 2373..2868
contig 04 2969..3469
contig 05 3570..3886
contig 06 3987..4673
contig 07 4774..5688
contig 08 5789..6264
contig 09 6365..6870
contig 10 6971..7382
contig 11 7483..7903
contig 12 8004..8333
contig 13 8434..8903
contig 14 9004..9420
contig 15 9521..9680
contig 16 9781..10953
contig 17 11094..12178
contig 18 12279..13325
contig 19 13426..14231
contig 20 14332..14718
contig 21 14819..15502
contig 22 15603..15740
contig 23 15841..16159
contig 24 16260..16507
contig 25 16608..17067
contig 26 17168..17424
contig 27 17525..17641
contig 28 17742..18174
contig 29 18275..18414
contig 30 18515..18723
contig 31 18824..19217
contig 32 19318..19460
contig 33 19561..19667
contig 34 19768..20117
contig 35 20218..20654
contig 36 20755..20894
contig 37 20985..21387
contig 38 21488..21774
contig 39 21875..22052
contig 40 22153..22547
contig 41 22648..22955
contig 42 23056..23543
contig 43 23644..23773
contig 44 23874..24139
contig 45

contig 46 24240..24602
contig 47 24703..25176
contig 48 25277..25673
contig 49 25774..26363
contig 50 26464..26897
contig 51 26998..29242
contig 52 29343..29710
contig 53 29811..29973
contig 54 30074..30247
contig 55 30348..31308
contig 56 31409..34048
contig 57 34149..35157
contig 58 35258..36473
contig 59 36574..37246
contig 60 37347..37850
contig 61 37951..38902
contig 62 39003..39823
contig 63 39924..40334
contig 64 40435..40887
contig 65 40988..41046
contig 66 41147..41464
contig 67 41565..42047
contig 68 42148..43221
contig 69 43322..44302
contig 70 44403..45038
contig 71 45139..45672
contig 72 45773..46403
contig 73 46504..47013
contig 74 47114..47546
contig 75 47647..47978
contig 76 48079..48523
contig 77 48624..49141
contig 78 49242..49811
contig 79 49912..51036
contig 80 51137..51679
contig 81 51780..52266
contig 82 52367..53022
contig 83 53123..53855
contig 84 53956..54255
contig 85 54356..54832
contig 86 54933..55626
contig 87 55727..56558
contig 88 56659..57428
contig 89 57529..58075
contig 90 58176..59033
contig 91 59134..59693
contig 92 59794..60166
contig 93 60267..60872
contig 94 60973..61737
contig 95 61838..62593
contig 96 62694..63459
contig 97 63560..64737
contig 98 64898..65488
contig 99 65589..67044
contig 100 67145..67584
contig 101 67685..68823
contig 102 68924..69802
contig 103 69903..70536
contig 104 70637..71214
contig 105 71315..71625
contig 106 71726..72513
contig 107 72614..74090
contig 108 74191..74968
contig 109 75069..76178
contig 110 76279..76723
contig 111 76824..78451
contig 112 78552..80226
contig 113 80327..81354
contig 114 81455..83535
contig 115 83636..85393
contig 116 85494..87577
contig 117 87678..90103
contig 118 90404..91635

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contig 119 91736..92784
 contig 120 92885..93370
 contig 121 93471..93972
 contig 122 94073..96680
 contig 123 96781..97314
 contig 124 97415..97765
 contig 125 97866..98251
 contig 126 98352..98406
 contig 127 98507..98635
 contig 128 98736..99395
 contig 129 99496..100370
 contig 130 100471..100855
 contig 131 100956..101534
 contig 132 101635..102482
 contig 133 102583..103329
 contig 134 103430..104188
 contig 135 104289..104450
 contig 136 104551..105037
 contig 137 105138..105707
 contig 138 105808..106278.
 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 138 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

1 753: contig of 753 bp in length
 754 853: gap of 100 bp
 854 1659: contig of 806 bp in length
 1660 1755: gap of 100 bp
 1760 2272: contig of 513 bp in length
 2273 2372: gap of 100 bp
 2373 2868: contig of 496 bp in length
 2869 3469: gap of 100 bp
 3470 3569: contig of 501 bp in length
 3570 3886: contig of 317 bp in length
 3887 4673: gap of 100 bp
 4674 4773: gap of 100 bp
 4774 5688: contig of 915 bp in length
 5689 5789: gap of 100 bp
 5790 6264: contig of 476 bp in length
 6265 6365: gap of 100 bp
 6366 6871: contig of 506 bp in length
 6872 7382: gap of 100 bp
 7383 7482: contig of 412 bp in length
 7483 7903: gap of 100 bp
 7904 8003: contig of 421 bp in length
 8004 8333: gap of 100 bp
 8334 8434: gap of 100 bp
 8435 8903: contig of 470 bp in length
 8904 9003: gap of 100 bp
 9004 9420: contig of 417 bp in length
 9421 9520: gap of 100 bp
 9521 9681: contig of 160 bp in length
 9682 10993: gap of 100 bp
 10994 10993: contig of 1213 bp in length

Query Match 79.1%; Score 17.4; DB 2; Length 106278;
 Best Local Similarity 94.7%; Pred. No. 2.4e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGAGC 22
 |||||
 Db 81934 CGAGCGGGCGGGCGGAGC 81916

RESULT 39
 AC132215

LOCUS AC132215 126323 bp DNA linear PLN 03-SEP-2002
 DEFINITION Genomic sequence for Oryza sativa, Nipponbare strain, clone
 OSUNBa0076506, from chromosome 3, complete sequence.
 ACCESSION AC132215
 VERSION AC132215.1 GI:22657512
 KEYWORDS HTG.
 SOURCE Oryza sativa (japonica cultivar-group)
 ORGANISM Oryza sativa (japonica cultivar-group)
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Ehrhartoideae; Oryzoideae; Oryza.
 REFERENCE 1 (bases 1 to 126323)
 AUTHORS McCombie, W.R., de la Bastide, M., Spiegel, L., Preston, R.,
 Nascentino, L., Zutavern, T., Ballia, V., Bell, M., Miller, B.,
 Katzenberger, F., Muller, S., Sullivan, P., Yang, C., Dike, S.,
 O'Shaughnessy, A., Palmer, L. and Dedhia, N.
 TITLE Genomic sequence for Oryza sativa, Nipponbare strain, clone
 OSUNBa0076506, from chromosome 3, complete sequence
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 126323)
 AUTHORS McCombie, W.R.
 TITLE Direct Submission
 JOURNAL Submitted (03-SEP-2002) Lita Annenberg Hazen Genome Center, Cold
 Spring Harbor Laboratories, 1, Bungtown Road, Cold Spring Harbor,
 NY 11724, USA

COMMENT This sequence was finished as follows unless otherwise noted: all
 regions were either double-stranded or sequenced with an alternate
 chemistry or covered by high quality data (i.e., phred quality >=
 30); an attempt was made to resolve all sequencing problems, such
 as compressions and repeats; all regions were covered by at least
 one plasmid subclone or more than one M13 subclone; and the
 assembly was confirmed by restriction digest. The nucleotide
 sequence of this BAC clone was generated by combining Syngenta,
 Monsanto and Cold Spring Harbor Laboratory Genome Center sequencing
 data.

FEATURES

Location/Qualifiers
 1..126323
 /organism="Oryza sativa (japonica cultivar-group)"
 /mol_type="genomic DNA"
 /cultivar="Nipponbare"
 /db_xref="taxon:39947"
 /chromosome="3"
 /clone="OSUNBa0076506"
 /clone_lib="HindIII"
 4490..4506
 misc_feature
 /note="We believe the assembly to be correct. The
 sequence is a mononucleotide (C) repeat in which the exact
 number of Cs is unknown. The majority of subclones in
 the area agree with the assembly, however, two subclones
 show two additional Cs."
 91540..91640
 misc_feature
 /note="We believe the assembly to be correct. The
 sequence is covered by a PCR product which was amplified
 with a high fidelity polymerase. The sequence is of high
 quality and there is partial coverage by several subclones
 with quality below phred30."
 35405 a 27507 c 26103 g 37308 t

BASE COUNT 35405 a 27507 c 26103 g 37308 t
 ORIGIN

Query Match 79.1%; Score 17.4; DB 8; Length 126323;
 Best Local Similarity 94.7%; Pred. No. 2.4e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 TCGATCGGGCGGGCGGAGC 21
 |||||
 Db 4300 TCGATCGGGCGGGCGGAGC 4318

RESULT 40
 AC137547
 LOCUS AC137547 144969 bp DNA linear PLN 02-APR-2003
 DEFINITION Oryza sativa (japonica cultivar-group) chromosome 3 clone
 OSUNBa0052J20, complete sequence.

AC137547
 VERSION AC137547.2 GI:29469501
 KEYWORDS HTG.
 SOURCE Oryza sativa (japonica cultivar-group)
 ORGANISM Oryza sativa (japonica cultivar-group)
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Ehrhartoideae; Oryzoideae; Oryza.
 REFERENCE 1 (bases 1 to 144969)
 AUTHORS Wing,R.A., Yu,Y., Soderlund,C., Kim,H.-R., Rambo,T., Currie,J. and
 Collura,K.
 TITLE Rice Genomic Sequence
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 144969)
 AUTHORS Wing,R.A., Yu,Y., Soderlund,C., Kim,H.-R., Rambo,T., Saski,C.,
 Currie,J. and Collura,K.
 TITLE Direct Submission
 JOURNAL Submitted (23-NOV-2002) Arizona Genomics Institute, University of
 Arizona, 303 Forbes, Tucson, AZ 85721, USA
 REFERENCE 3 (bases 1 to 144969)
 AUTHORS Wing,R.A., Yu,Y., Soderlund,C., Kim,H.-R., Rambo,T., Currie,J. and
 Collura,K.
 TITLE Direct Submission
 JOURNAL Submitted (02-APR-2003) Arizona Genomics Institute, University of
 Arizona, 303 Forbes, Tucson, AZ 85721, USA
 COMMENT On Apr 2, 2003 this sequence version replaced gi:25189927.
 This sequence was finished as follows unless otherwise noted: all
 regions were either double-stranded or sequenced with an alternate
 chemistry or covered by high quality data (i.e., phred quality
 >30); an attempt was made to resolve all sequencing problems, such
 as compressions and repeats; all regions were covered by more than
 one plasmid subclone; and the assembly was confirmed by
 restriction digest. There is a TA dinucleotide repeat ranging from
 10-30 pairs at 92656-91715. From base 10472-10575 the subclone reads
 for OSJNBa0052J20 have deleted sequence but the PCR done off this
 BAC DNA matches the overlapping clone OSJNBb0014110. There are
 Bacterial Transposons at the following locations: 28628-33998,
 45024-48063 and 43861-47164. At bases 44656-44749 only transposon
 reads cover this area. There is a repeat of TAA (36-38 repeats) at
 base 50810. The assembly overlaps from base 1-33264 with
 OSJNBb0014110 (accession #AC12622). The overlap is from bases
 80683-113946 on OSJNBb0014110. The assembly overlaps from base
 127973-144969 with OSJNBa0076E06 (accession #AC132215). The overlap
 is from bases 1-15998 on OSJNBa0076E06. The nucleotide sequence of
 this BAC clone was generated by combining Syngenta, Monsanto and
 Arizona Genomics Institute sequencing data.

FEATURES

Location/Qualifiers
 1..144969
 source
 /organism="Oryza sativa (japonica cultivar-group)"
 /mol_type="genomic DNA"
 /db_xref="taxon:39947"
 /chromosome="3"
 /clone="OSJNBa0052J20"

BASE COUNT 40242 a 32367 c 31769 g 40591 t
 ORIGIN

Query Match 79.1%; Score 17.4; DB 8; Length 144969;
 Best Local Similarity 94.7%; Pred. No. 2.4e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAG 21

Db 132271 TCGATCGGGCGGGCGGG 132289
 |||||

Search completed: February 18, 2004, 16:42:29
 Job time : 1543 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 16:09:36 ; Search time 172 Seconds
(without alignments)
345.277 Million cell updates/sec

Title: US-10-026-341A-2

Perfect score: 22

Sequence: 1 attgatcgggcgggcgagc 22

Scoring table: IDENTITY_NUC

Gapop 10_0 , Gapext 1.0

Searched: 2552756 seqs, 1349719017 residues

Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

N_Geneseq_19Jun03:*

- 1: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1980.DAT:*
- 2: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1981.DAT:*
- 3: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1982.DAT:*
- 4: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1983.DAT:*
- 5: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1984.DAT:*
- 6: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1985.DAT:*
- 7: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1986.DAT:*
- 8: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1987.DAT:*
- 9: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1988.DAT:*
- 10: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1989.DAT:*
- 11: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1990.DAT:*
- 12: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1991.DAT:*
- 13: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1992.DAT:*
- 14: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1993.DAT:*
- 15: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1994.DAT:*
- 16: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1995.DAT:*
- 17: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1996.DAT:*
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- 19: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1998.DAT:*
- 20: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA1999.DAT:*
- 21: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA2000.DAT:*
- 22: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA2001A.DAT:*
- 23: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA2001B.DAT:*
- 24: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA2002.DAT:*
- 25: /SIDSL/gcgdata/geneseq/geneseq-nbml/NA2003.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Query | Score | Match | Length | ID | Description |
|------------|-------|-------|-------|--------|----------|--------------------|
| 1 | 22 | 100.0 | 22 | 17 | AAT18820 | SPI motif. Homo s |
| 2 | 22 | 100.0 | 22 | 18 | AAT77128 | Spi consensus. Sy |
| 3 | 22 | 100.0 | 22 | 20 | AAX76050 | CAMP response elem |
| 4 | 22 | 100.0 | 22 | 21 | AAD01054 | Oligonucleotide wi |
| 5 | 22 | 100.0 | 22 | 21 | AAA46366 | Nucleotide sequenc |
| 6 | 22 | 100.0 | 22 | 22 | AAD10308 | SPI oligonucleotid |
| 7 | 22 | 100.0 | 22 | 22 | AAD06435 | PCR primer Spi, us |
| 8 | 22 | 100.0 | 22 | 24 | AAD44321 | Decoy Spi oligonuc |

| | | | | | | |
|----|------|-------|-------|----|----------|--------------------|
| 9 | 22 | 100.0 | 22 | 24 | AAD40710 | Mouse osterix bind |
| 10 | 22 | 100.0 | 22 | 24 | ABQ74742 | Glucose carrier ty |
| 11 | 22 | 100.0 | 22 | 24 | ABA92271 | Sp-1 wild-type oli |
| 12 | 22 | 100.0 | 22 | 24 | ABA93509 | Regulatory element |
| 13 | 22 | 100.0 | 22 | 25 | ABZ58136 | Transcription fact |
| 14 | 22 | 100.0 | 49 | 22 | AAF61841 | SPI-specific DNA p |
| 15 | 22 | 100.0 | 49 | 22 | AAF61841 | SPI-specific DNA p |
| 16 | 22 | 100.0 | 49 | 22 | AAF61843 | SPI-specific DNA p |
| 17 | 22 | 100.0 | 49 | 22 | AAF61843 | SPI-specific DNA p |
| 18 | 21 | 95.5 | 21 | 22 | AAF99620 | Immunostimulatory |
| 19 | 21 | 95.5 | 21 | 22 | AAF99621 | Immunostimulatory |
| 20 | 21 | 95.5 | 21 | 24 | ABS78341 | Angiogenesis inhib |
| 21 | 21 | 95.5 | 21 | 24 | ABS78342 | Angiogenesis inhib |
| 22 | 21 | 95.5 | 21 | 24 | ABL38746 | Immunostimulatory |
| 23 | 21 | 95.5 | 21 | 24 | ABL38839 | Immunostimulatory |
| 24 | 21 | 95.5 | 21 | 25 | ABT17254 | Transcription fact |
| 25 | 21 | 95.5 | 21 | 25 | ABT17255 | Transcription fact |
| 26 | 21 | 95.5 | 63 | 25 | ABT17322 | Transcription fact |
| 27 | 20.4 | 92.7 | 22 | 24 | ABX89728 | Oestrogen response |
| 28 | 20 | 90.9 | 20 | 19 | AAV46004 | Immune adjuvant sp |
| 29 | 20 | 90.9 | 20 | 21 | AAZ99627 | Nucleotide sequenc |
| 30 | 20 | 90.9 | 20 | 21 | AAZ99627 | Murine Toll-like r |
| 31 | 18.8 | 85.5 | 22 | 24 | ABL39229 | Sp-1 mutant oligon |
| 32 | 18.4 | 83.6 | 22 | 15 | AAQ67304 | Detection probe fo |
| 33 | 18 | 81.8 | 45 | 13 | AAQ30483 | Oligonucleotide co |
| 34 | 18 | 81.8 | 45 | 13 | AAQ30483 | Oligonucleotide co |
| 35 | 18 | 81.8 | 47 | 13 | AAQ30484 | Oligonucleotide co |
| 36 | 17.4 | 79.1 | 183 | 25 | ABA00746 | Enhl enhancer. Sy |
| 37 | 17.4 | 79.1 | 722 | 24 | ABQ44940 | Oligonucleotide fo |
| 38 | 17.4 | 79.1 | 722 | 24 | ABQ44941 | Oligonucleotide fo |
| 39 | 17.4 | 79.1 | 747 | 24 | ABQ43520 | Oligonucleotide fo |
| 40 | 17.4 | 79.1 | 747 | 24 | ABQ43521 | Oligonucleotide fo |
| 41 | 17.4 | 79.1 | 9646 | 24 | ABL33688 | Human immune syste |
| 42 | 17.2 | 78.2 | 10035 | 25 | ABZ66813 | Orthosomycin biosy |
| 43 | 17.2 | 78.2 | 11115 | 23 | ABL50562 | Micronospora ear |
| 44 | 17 | 77.3 | 540 | 24 | ABQ23090 | Oligonucleotide fo |
| 45 | 17 | 77.3 | 540 | 24 | ABQ23091 | Oligonucleotide fo |

ALIGNMENTS

RESULT 1

AAT18820
ID AAT18820 standard; DNA; 22 BP.
XX AAT18820;
XX AC
XX 17-AUG-1996 (first entry)
DT
XX
DE SPI motif.
XX
XX NF-X1; transcription factor; major histocompatibility complex; MHC;
XX allergy; HLA-DRA; ds.
XX
OS Homo sapiens.
XX
XX WO9612823-A1.
XX
XX 02-MAY-1996.
XX
XX 20-OCT-1995; 95WO-US12749.
XX
XX 21-OCT-1994; 94US-0327832.
XX
XX (HARD) HARVARD COLLEGE.
XX (UYJO) UNIV JOHNS HOPKINS.
XX
XX Ono SJ, Strominger JL;
XX WPI; 1996-230621/23.
XX
XX Transcription factor, NF-X1 and DNA encoding it - used in regulation
PT

PT of MHC class II expression and in treatment of allergic disease

XX Example 4; Page 42; 93pp; English.

CC Recombinant transcription factor NF-X1 (see AAR94957) forms a
 CC specific complex with the HLA-DRA X1 box oligonucleotide (AAT18817)
 CC which is competed for by 100-fold excess cold, double-stranded
 CC oligonucleotides containing the analogous regions from the HLA-DRB,
 CC -DPA, -DPB, -DQA and -DQB promoters, but not by HLA-DRA Y-box
 CC (AAT18818), S-box (AAT18819), S1 (AAT18820) or the interferon-beta gene
 CC positive-regulatory domain II element (AAT18821). It is concluded
 CC that NF-X1 binds sequence-specifically with all human class II
 CC major histocompatibility X1 boxes (see also AAT18812).

XX SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 100.0%; Score 22; DB 17; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22

Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 2

AAT77128
 ID AAT77128 standard; DNA; 22 BP.

AC AAT77128;

DT 07-DEC-1997 (first entry)

DE S1 consensus.

XX 17-Beta-hydroxysteroid dehydrogenase type I; HSD17B1; human;
 KW promoter; ds.

XX Synthetic.

XX WO9720942-A1.

PN 12-JUN-1997.

PD 04-DEC-1996; 96WO-FI00647.

XX 05-DEC-1995; 95US-0007976.

XX (OIK/) OIKARINEN J A.

PA (BELT/) PELTOKETO E H.

PA (PIAO/) PIAO Y.

PA (VIHK/) VIHKO R K.

XX Oikarinen JA, Peltoketo EH, Piao Y, Viikko RK;

XX WPI; 1997-319789/29.

XX Human 17-beta-hydroxysteroid dehydrogenase type I (HSD17B1)
 PT transcription regulatory elements - used for identifying agents
 PT which can up- or down-regulate HSD17B1 expression to increase or
 PT decrease oestrogen production

XX Example 6; Page 38; 69pp; English.

XX This oligonucleotide comprises a consensus sequence for S1
 CC binding sites. It was used with oligonucleotides (see
 CC AAT77122-24 and AAT77126-27) based on the promoter region of the human
 CC 17-beta-hydroxysteroid dehydrogenase type I (HSD17B1) gene (see
 CC AAT77112), and with an AP-2 consensus oligonucleotide (see AAT77125),
 CC in the detailed characterisation of the HSD17B1 promoter, and to
 CC examine the role of S1 and AP-2 binding sites in promoter
 CC function.

XX

SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 100.0%; Score 22; DB 18; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22

Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 3

AA76050
 ID AA76050 standard; DNA; 22 BP.

AC AA76050;

DT 30-JUL-1999 (first entry)

DE CAMP response element oligonucleotide SEQ ID NO:18.

XX CRE; cAMP response element; transcription factor decoy; cis-element;
 KW tumour growth inhibitor; palindromic; hairpin; cancer; metabolism;
 KW gene transcription regulation; inhibiting proliferation; ds.

XX Synthetic.

XX WO9926634-A1.

XX 03-JUN-1999.

PF 23-NOV-1998; 98WO-US25307.

XX 24-NOV-1997; 97US-0977643.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Cho-Chung YS;

XX WPI; 1999-347612/29.

XX Nucleic acids that compete with response elements for transcription
 PT factors

XX Example 10; Page 54; 83pp; English.

XX The present invention describes a composition (A) comprising one or more
 CC nucleic acids (I) that compete with cAMP (cyclic adenosine monophosphate)
 CC response element (CRE) enhancer DNA for binding to transcription factors
 CC (TF). (I) are used to regulate gene transcription in cells, in vitro or
 CC in vivo, specifically for inhibiting proliferation of cancer cells, but
 CC possibly also for regulation of metabolism in hepatitis B and other
 CC viruses. HCT-15 human multidrug resistant colon carcinoma cells (2
 CC million) were inoculated subcutaneously into the flank of nude mice,
 CC then the CRE oligonucleotide 5'-TGACGTTTCATGAGTTTCATCA-3' injected
 CC intraperitoneally at doses of 0.1 mg, 5 times per week, once the tumour
 CC had reached 30-50 mg. This treatment resulted in over 85% reduction in
 CC tumour growth, relative to an untreated control. (I) have high affinity
 CC for TF and can inhibit growth of cancer cells without adverse effects on
 CC normal cells (contrast use of antisense RNA). The method does not
 CC require knowledge of the target gene sequence, only of the response
 CC element sequence. The present sequence is used in the exemplification
 CC of the present invention.

XX SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 100.0%; Score 22; DB 20; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22

Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 4
AAD01054
ID AAD01054 standard; DNA; 22 BP.
XX
AC AAD01054;
XX
DT 04-OCT-2000 (first entry)
XX
DE Oligonucleotide with Spl consensus site for EMSA of human cyclin A1 gene.
XX
KW Human; cyclin A1; transgenic; stem cell; male germ cell; promoter;
KW fluorescent protein; light emitting reporter protein; genetic defect;
KW gene therapy; molecular engineering; trauma; vertebrate; Spl;
KW electrophoretic mobility shift assay; EMSA; ds.
XX
OS Unidentified.
XX
PN WO200029602-A1.
XX
PD 25-MAY-2000.
XX
PF 15-APR-1999; 99WO-US08277.
XX
PR 13-NOV-1998; 98US-0191920.
XX
PR 13-NOV-1998; 98WO-US24238.
XX
PA (CEDA-) CEDARS SINAI MEDICAL CENT.
PA (UNLO) IMPERIAL COLLEGE SCI TECHNOLOGY & MED.
XX
PI Readhead CW, Winston R, Koeffler HP, Muller C;
XX
DR WPI; 2000-387812/33.
XX
PT Producing selectable transgenic stem cells for use in molecular
PT engineering, e.g. creating transgenic animals -
XX
PS Example 12; Page 41; 98pp; English.
XX
CC The patent discloses in vivo and in vitro methods for producing
CC selectable transgenic stem cells by transfecting male germ cells with a
CC transfection mixture comprising a nucleic acid construct containing a
CC transcriptional unit of a stem cell-specific promoter, e.g. cyclin
CC A1 promoter, operably linked to a gene encoding a fluorescent or
CC light emitting protein. These methods may be used for producing
CC transgenic vertebrate animals and for repairing genetic defects.
CC They are suitable for germ line and stem cell line gene therapy.
CC They are also valuable in identifying cell lineages before full
CC differentiation and to facilitate modification and/or engineering of
CC specific tissues in vitro for their subsequent transplantation in the
CC treatment of disease or trauma. The present sequence is an
CC oligonucleotide, containing Spl consensus site, used in electrophoretic
CC mobility shift assay using nuclear extract from Hela cells to
CC analyse human cyclin A1 gene. Cyclin A1 promoter derived from
CC the 5' end of cyclin A1 gene is stem cell-specific and used in a
CC DNA construct for producing selectable transgenic stem cells.
XX
SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
Query Match 100.0%; Score 22; DB 21; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 5
AAA46366/c
ID AAA46366 standard; DNA; 22 BP.
XX

AC AAA46366;
XX
DT 04-SEP-2000 (first entry)
XX
DE Nucleotide sequence of a Spl element oligonucleotide.
XX
KW Malignant cell; OP2 C2-6; OP7 C3-1; OP9 A4-2; OP11 C1-3; OP11 G2-10;
KW FAS OP13; FAS OP17; CINN 1; CINN 2; anti-neoplastic drug; malignancy;
KW cancer; brain cancer; intracranial neoplasm; ss.
XX
OS Unidentified.
XX
PN WO200028090-A2.
XX
PD 18-MAY-2000.
XX
PF 12-NOV-1999; 99WO-US27005.
XX
PR 12-NOV-1998; 98US-0108120.
XX
PR 27-JUL-1999; 98US-0145640.
XX
PA (NYXI-) NYXIS INC.
XX
PI Kroes RA, Moskal JR, Jastrow A, Yamamoto H;
XX
DR WPI; 2000-376595/32.
XX
PT Method for ascertaining the propensity of a cell for malignant
PT phenotype, used for diagnosis of cancers, including those of the lung,
PT breast and brain -
XX
PS Disclosure; Page 44; 66pp; English.
XX
CC The specification describes a method for ascertaining the propensity of
CC a cell for malignant phenotype. The method comprises assaying a cell or
CC biological sample for a signal indicating the transcription of a nucleic
CC acid transcript. The transcript is from at least one gene selected from
CC the genes encoded for by or containing the characteristic nucleic acid
CC sequences of OP2 C2-6, OP7 C3-1, OP9 A4-2, OP11 C1-3, OP11 G2-10,
CC FAS OP13, FAS OP17, CINN 1 or CINN 2 (AAA46354-62, respectively). Cells
CC identified using the method can be used for ascertaining the suitability
CC of an anti-neoplastic drug candidate for efficacy in treating a
CC malignancy. The method can be used in the diagnosis of many types of
CC cancers, including cancers of the breast, prostate, colon and lung. It
CC may also be used for diagnosis and treatment of brain cancers, including
CC intracranial neoplasms such as those of the skull, meninges, cranial
CC nerves, neuroglia, ependyma, pituitary or pineal body, as well as those
CC of metastatic origin. The present sequence represents a Spl element
CC oligonucleotide, which is used in the course of the invention.
XX
SQ Sequence 22 BP; 3 A; 11 C; 5 G; 3 T; 0 other;
Query Match 100.0%; Score 22; DB 21; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 22 ATTCGATCGGGCGGGCGGAGC 1

RESULT 6
AAD10308
ID AAD10308 standard; DNA; 22 BP.
XX
AC AAD10308;
XX
DT 24-SEP-2001 (first entry)
XX
DE Spl oligonucleotide to determine nuclear factor kappa B activation.
XX
KW Human; Par-4; presenilin; PS; neuroprotective; nuclear factor kappa B;
KW NF-kappa B; neuronal degeneration; spinal muscular atrophy; paralysis;

KW peripheral neuropathy; motorneuron disorder; neurodegenerative disorder;
KW Parkinson's disease; Meniere's disease; multiple sclerosis; Bell's palsy;
KW Huntington's chorea; Down's syndrome; amyotrophic lateral sclerosis; ALS;
KW nerve deafness; Alzheimer's disease; epilepsy; ds.
OS Unidentified.
FN WO200151671-A2.
XX 19-JUL-2001.
XX 08-JAN-2001; 2001WO-US00526.
XX 10-JAN-2000; 2000US-0175200.
PR 04-JAN-2001; 2001US-0754949.
XX (SCIO-) SCIOS INC.
XX McCarthy J, Cordell B;
PI WPI; 2001-451872/48.
XX
XX Identifying inhibitors of neuronal degeneration useful for treating
PT e.g. Alzheimer's disease, by determining the ability of a compound to
PT induce nuclear factor kappa B activation, with the involvement of
PT presenilin or Par-4 -
XX Example 5; Page 25; 66pp; English.
PS The invention relates to human Par-4 protein, presenilin protein (PS1
XX and PS2) and their corresponding DNA molecules. The invention also
CC relates to a method for identifying inhibitors of neuronal degeneration,
CC comprising cotransfecting eukaryotic host cells expressing presenilin
CC (PS), with a Par-4 DNA, and an NF-kappa B dependent reporter construct,
CC exposing the cotransfected cells to a candidate molecule and monitoring
CC the ability of the candidate molecule to induce NF-kappa B activation.
CC Presenilin proteins participate in nuclear factor kappa B (NF-kappa B)
CC signalling and activation. The inhibitors of neuronal degeneration
CC are useful for treating neurodegenerative disorders such as Alzheimer's
CC disease, Parkinson's disease, epilepsy, multiple sclerosis, Huntington's
CC chorea, Down's syndrome, nerve deafness, Meniere's disease and also for
CC treating peripheral neuropathies, motorneuron disorders such as
CC amyotrophic lateral sclerosis (ALS), Bell's palsy and various conditions
CC involving spinal muscular atrophy and paralysis. The present DNA sequence
CC is SP-1 oligonucleotide which is used to determine nuclear factor kappa B
CC (NF-kappaB) activation.
XX
XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
SQ Query Match 100.0%; Score 22; DB 22; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTCGATCGGGGGGGCGGCGAGC 22
Db 1 ATTCGATCGGGGGGGCGGCGAGC 22
RESULT 7
ID AAD06435 standard; DNA; 22 BP.
XX AAD06435;
XX 10-AUG-2001 (first entry)
DT PCR primer Sp1, used in electrophoretic mobility shift assay.
DE Human; Mcl-1 gene regulatory element; Mcl-1s/deltaTM variant;
XX neuronal cell; tumour cell; apoptosis; therapy; cancer; psoriasis;
KW diabetic retinopathy; corneal graft neovascularisation;
KW neovascular glaucoma; epithelial condition; autoimmune disease;
KW rheumatoid arthritis; systemic lupus erythematosus;

KW neurodegenerative disease; PCR primer; ss.
XX Homo sapiens.
XX WO200136594-A1.
PD 25-MAY-2001.
XX 14-JAN-2000; 2000WO-US00969.
PF 16-NOV-1999; 99US-0166113.
XX (DART-) DARTMOUTH COLLEGE.
PA Craig RW, Bingle CD, Whyte M;
PI WPI; 2001-343812/36.
XX Novel Mcl-1 gene regulatory elements, useful for modulating expression
PT of Mcl-1 polypeptide or its variant which regulate apoptosis in
PT neuronal or tumor cells -
XX Example 1; Page 63; 125pp; English.
PS The present invention relates to Mcl-1 gene regulatory elements and the
XX variant Mcl-1s/deltaTM. The anti-apoptotic Mcl-1 protein is encoded by
CC exons 1, 2 and 3. The pro-apoptotic Mcl-1s/deltaTM variant encoded by
CC exons 1 and 3 is obtained due to alternative mRNA splicing. The Mcl-1
CC gene regulatory element is useful for modulating the Mcl-1 gene
CC expression in a cell e.g., neuronal cell or tumour cell, such that
CC apoptosis of the cell is induced or cell viability is increased. The
CC Mcl-1 and its regulatory elements are used for treating pathological
CC conditions which include cancer, diabetic retinopathy, corneal graft
CC neovascularisation and neovascular glaucoma, epithelial conditions such
CC as psoriasis, autoimmune diseases like rheumatoid arthritis, systemic
CC lupus erythematosus, and neurodegenerative diseases. The present sequence
CC is a PCR primer Sp1, used in the electrophoretic mobility shift assay,
CC which is related to the invention.
XX
XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
SQ Query Match 100.0%; Score 22; DB 22; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTCGATCGGGGGGGCGGCGAGC 22
Db 1 ATTCGATCGGGGGGGCGGCGAGC 22
RESULT 8
ID AAD44321 standard; DNA; 22 BP.
XX AAD44321;
XX 13-DEC-2002 (first entry)
DT Decoy Sp1 oligonucleotide.
DE Fibrotic condition; gene expression; cirrhosis; hypertrophic scar;
XX gene therapy; fibrosis; skin disorder; sclerodermic lesion; keloid;
KW trauma; surgery; Sp1; ds.
XX Unidentified.
XX WO200266071-A2.
FN 29-AUG-2002.
XX 21-DEC-2001; 2001WO-US49141.
XX 03-JAN-2001; 2001US-259585P.
PR

XX PA (UYJE-) UNIV JEFFERSON THOMAS.
 XX PI Mauviel A;
 XX DR WPI; 2002-667041/71.
 XX PT Treating a fibrotic condition, e.g. cirrhosis, comprises administering
 XX PT antisense Spl or decoy Spl oligonucleotides that inhibit transcription
 XX PT or gene expression of an extra-cellular matrix gene or transforming
 XX PT growth factor-beta.
 XX XX
 XX PS Claim 5; Page 38; 38pp; English.
 XX XX
 XX CC The invention relates to a method of treating a fibrotic condition (the
 XX CC transcription or gene expression of an extra-cellular matrix (ECM) gene
 XX CC or transforming growth factor-beta (TGF-beta) is inhibited in a mammal).
 XX CC The method involves administering an antisense Spl or a decoy Spl
 XX CC oligonucleotide. The method, antisense Spl and decoy Spl oligonucleotide
 XX CC are useful for treating fibrotic conditions e.g. cirrhosis, radiation
 XX CC induced fibrosis, skin disorders (sclerodermic lesions, hypertrophic
 XX CC scars, keloids), kidney fibrosis, lung fibrosis and myelofibrosis. They
 XX CC are also useful for treating or preventing fibrotic scarring following
 XX CC trauma or surgery. The invention is useful in gene therapy. The present
 XX CC sequence is decoy Spl oligonucleotide..
 XX XX
 XX SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
 Query Match 100.0%; Score 22; DB 24; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
 DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 9
 ID AAD40710 standard; DNA; 22 BP.
 XX AC AAD40710;
 XX DT 30-OCT-2002 (first entry)
 XX DE Mouse osterix binding oligonucleotide, Spl.
 XX KW Bone formation; transcription factor; osteoblast; Paget's disease;
 XX KW osterix protein; glucocorticoid; osteoporosis; cytokine;
 XX KW periodontal disease; tooth loss; bone fracture; rheumatoid arthritis;
 XX KW metastatic bone disease; gene therapy; growth factor; osteopathic;
 XX KW differentiation; mouse; ss.
 XX OS Mus musculus.
 XX FN WO200244380-A2.
 XX PD 06-JUN-2002.
 XX XX
 XX PF 30-NOV-2001; 2001WO-US44898.
 XX PR 30-NOV-2000; 2000US-0734329.
 XX PA (TEXA) UNIV TEXAS SYSTEM.
 XX PI De Crombrughe B, Nakashima K, Zhou X;
 XX PS WPI; 2002-519587/55.
 XX XX
 XX CC Novel DNA segment encoding Osterix polypeptide which is a master
 XX CC transcription factor that controls osteoblast differentiation and is
 XX CC useful for treating osteoporosis, in patient by stimulating bone
 XX CC formation.

XX PS Example 4; Page 107; 144pp; English.
 XX XX
 XX CC The invention relates to a master bone formation transcription factor
 XX CC that controls osteoblast differentiation, osterix protein and its
 XX CC corresponding nucleic acid sequence. Osterix protein and its DNA and
 XX CC agents that interact with the protein to activate or stimulate the
 XX CC differentiation of bone cells is used for treatment of glucocorticoid
 XX CC induced osteoporosis, Paget's disease, periodontal disease, tooth loss,
 XX CC bone fractures, rheumatoid arthritis, metastatic bone disease, etc.
 XX CC Osterix DNA is useful as probes or primers in nucleic acid hybridisation
 XX CC experiments. It is also used in gene therapy. Osterix protein is useful
 XX CC for controlling bone formation, serves as receptors of soluble
 XX CC molecules, e.g. cytokines, growth factors, etc., as homing/adhesion/
 XX CC rolling receptors mediating the migration of osteoblast, as signalling
 XX CC receptors, thereby regulating function of osteoblasts, and/or ligands
 XX CC of signalling receptors or osteoblast. It is also useful for purifying
 XX CC osteoblasts, from a test composition suspected of containing the cells.
 XX CC The present sequence is mouse osterix gene binding oligonucleotide used
 XX CC in the exemplification of the invention.
 XX XX
 XX SQ Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
 Query Match 100.0%; Score 22; DB 24; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
 DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 10
 ID ABQ74742 standard; DNA; 22 BP.
 XX AC ABQ74742;
 XX DT 24-OCT-2002 (first entry)
 XX DE Glucose carrier type 2 promoter related oligonucleotide #8.
 XX KW Hepatocyte nuclear factor; HNF; gene therapy; liver; pancreas;
 XX KW glucose carrier type 2 promoter; tissue-specific expression;
 XX KW glucose metabolism related disease; diabetes; ss.
 XX OS Synthetic.
 XX XX
 XX PN K32001109882-A.
 XX PD 12-DEC-2001.
 XX XX
 XX PF 03-JUN-2000; 2000KR-0030606.
 XX PR 03-JUN-2000; 2000KR-0030606.
 XX PA (GENE-) GENEPIA CO LTD.
 XX XX
 XX PI Ahn YH, Cha JY, Huh MU, Kim GS, Kim HI;
 XX PS WPI; 2002-398926/43.
 XX XX
 XX PT DNA sequence for tissue-specific expression in liver or pancreas,
 XX PT useful for gene therapy of glucose metabolism related disease, e.g.
 XX PT diabetes -
 XX XX
 XX PS Example; Page 6; 15pp; Korean.
 XX XX
 XX CC The present invention describes a liver or pancreas tissue-specific
 XX CC expression related DNA sequence (I). (I) is particularly a hepatocyte
 XX CC nuclear factor (HNF) binding region present in glucose carrier type 2
 XX CC promoter. (i) participates in tissue-specific expression in liver or
 XX CC pancreas and HNF1 and HNF3 characteristically bind to it. (i) can be

CC used in a recombinant vector useful for gene therapy in glucose
 CC metabolism related diseases, such as diabetes. The present sequence
 CC represents an oligonucleotide which is used in the exemplification of
 CC the present invention.

CC Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
 CC
 CC Query Match 100.0%; Score 22; DB 24; Length 22;
 CC Best Local Similarity 100.0%; Pred. No. 3.3;
 CC Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAGC 22
 |||||
 Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 11
 ABA92271
 ID ABA92271 standard; DNA; 22 BP.

XX AC ABA92271;
 XX
 XX
 XX 10-JUN-2002 (first entry)

DE Sp-1 wild-type oligonucleotide, used in EMSA.

XX SP-1; neuron; antialzheimers; antiparkinsonian; antisclerotic;
 KW neuroprotective; neurotropic; anticonvulsant; vascular; hypotensive;
 KW cerebroprotective; virucide; anti-HIV; diagnosis; therapy;
 KW electrophoretic mobility shift assay; EMSA; ds.

XX Homo sapiens.

XX WO200215912-A1.

XX 28-FEB-2002.

XX 24-AUG-2001; 2001WO-US26527.

XX 25-AUG-2000; 2000US-228201P.

XX 26-OCT-2000; 2000US-243295P.

XX (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.

XX Ratan RR, Chatterjee S;

XX WPI; 2002-242023/29.

XX Diagnosing and treating diseases associated with oxidative stress, DNA
 PT damage or growth factor depletion, e.g. Alzheimer's and Parkinson's, by
 PT administering e.g. mithramycin, chromomycin, daunomycin, olivomycin and
 PT WP631 -

XX Example 1; Page 23; 69pp; English.

XX The present sequence is that of a double-stranded wild-type Sp-1
 CC oligonucleotide, which was used in an electrophoretic mobility
 CC shift assay to determine the effect of oxidative stress on Sp-1 DNA
 CC binding, and to determine the effects of candidate compounds on Sp-1
 CC protein levels. Sp-1 DNA binding activity in cortical neurons was
 CC shown to be low, but was dramatically enhanced by oxidative stress.
 CC The invention provides methods for detecting and treating diseases
 CC associated with oxidative stress, DNA damage or growth factor
 CC depletion, and identifying agents for the treatment of such
 CC conditions. A compound is deemed to be an inhibitor of oxidative
 CC stress, DNA damage, growth factor depletion or cell death if it
 CC reduces the protein level of an Sp family member or if it decreases
 CC the binding of an Sp family member to DNA. A method for preventing
 CC or treating a disease or disorder of the nervous system, the ageing
 CC process or associated with apoptosis involves administering a
 CC compound that inhibits the induction of an Sp family member or the
 CC binding of an Sp family member to DNA, e.g. mithramycin, and
 CC chromomycin, daunomycin, olivomycin or WP631. Diseases and

CC disorders that can be treated include Alzheimer's disease,
 CC Creutzfeldt-Jacob disease, kuru, Huntington's disease, aneurysm,
 CC stroke associated with an increase in blood pressure, spinal cord
 CC disease, spinal cord injury, brain injury, multiple system atrophy,
 CC amyotrophic lateral sclerosis, progressive supranuclear palsy,
 CC neurodegeneration associated with the ageing process, mitochondrial
 CC disease, HIV infection, herpes infection and multiple sclerosis
 CC (all claimed).

XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 100.0%; Score 22; DB 24; Length 22;
 Best Local Similarity 100.0%; Pred. No. 3.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAGC 22
 |||||
 Db 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 12
 ABA93509

ID ABA93509 standard; DNA; 22 BP.

XX AC ABA93509;

XX 24-APR-2002 (first entry)

XX Regulatory element related oligonucleotide probe SEQ ID NO:9.

XX Pancreatic langerhans beta cell proliferation promoter; glucocorticoid;
 KW apoptosis inhibitor; regulatory element; HGF; hepatocyte growth factor;
 KW antidiabetic; apoptotic; diabetes; probe; ss.

XX Synthetic.

XX WO200193899-A1.

XX 13-DEC-2001.

XX 01-JUN-2001; 2001WO-JP04660.

XX 02-JUN-2000; 2000JP-0170447.

XX 28-FEB-2001; 2001JP-0054072.

XX (OKAM/) OKAMOTO H.

XX Okamoto H;

XX WPI; 2002-147646/19.

XX Promoters for pancreatic langerhans beta cell proliferation and
 PT apoptosis inhibitors, useful in efficiently screening candidate
 PT compounds for expression inducers of the genes, e.g. for treatment of
 PT diabetes -

XX Example 3; Page 30; 73pp; Japanese.

XX The present invention describes an expression inducer (I) of hepatocyte
 CC growth factor (HGF) gene and/or Reg gene containing a cytokine with
 CC gp130 as receptor and glucocorticoid. Also described are: (1) a promoter
 CC for pancreatic langerhans beta cell proliferation containing a cytokine
 CC with gp130 as receptor and glucocorticoid; (2) an apoptosis inhibitor
 CC containing a cytokine with gp130 as receptor and glucocorticoid; (3) a
 CC promoter for pancreatic langerhans beta cells containing HGF or a nucleic
 CC acid encoding HGF; (4) an apoptosis inhibitor of pancreatic langerhans
 CC beta cell proliferation containing HGF or a nucleic acid encoding HGF;
 CC (5) drugs containing Reg protein or a nucleic acid encoding Reg protein;
 CC (6) a method for screening candidate compounds as expression inducer of
 CC Reg protein comprising: (a) supplying cells transferred with a vector
 CC containing a reporter gene ligated functionally to the 5'-upstream of
 CC Reg gene carrying poly(ADP-ribose) synthase/polymerase (PARP) binding
 CC sequence; (b) contacting the cells with a test sample and detecting

CC reporter activity of the cells; and (c) selecting a compound that can
CC increase the reporter activity as compared to a control; (7) a method
CC for screening candidate compounds as expression inducers of HGF gene
CC comprising: (a) supplying cells introduced with a vector containing a
CC reporter gene ligated functionally to 5' upstream of HGF gene carrying a
CC 11-6/dexamethasone-responsive sequence; (b) contacting the cells with a
CC test sample and detecting reporter activity of the cells; and (c)
CC selecting a compound that can increase the reporter activity as compared
CC to a control. (i) has antidiabetic and apoptotic activities. The
CC promoters, inhibitors and screened compounds are for treating diabetes.
CC The present sequence represents a regulatory element related probe
CC which is used in an example from the present invention.

XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;
SQ Query Match 100.0%; Score 22; DB 24; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 13

ABZ58136
ID ABZ58136 standard; DNA; 22 BP.

XX AC ABZ58136;
XX XX

DT 22-APR-2003 (first entry)

DE Transcription factor Sp-1 probe.

XX Human; osteocalcin; promoter; bone; tumour; prostate cancer;
KW metastasis; gene therapy; diagnosis; prognosis; marker; cytostatic;
KW transcription factor; Sp-1; probe; ss.

OS Homo sapiens.

XX WO2003006621-A2.

PN 23-JAN-2003.

PD 12-JUL-2002; 2002WO-US222216.

PF 13-JUL-2001; 2001US-305360P.

PR (UYVI-) UNIV VIRGINIA PATENT FOUND.

PA Chung LWK, Yeung F;
PI WPI; 2003-221733/21.

DR Nucleic acid sequence for diagnosing, prognosing or treating calcified
PT tumors and tissues, e.g. prostate cancer, comprises an improved
PT recombinant human osteocalcin promoter activity -

PS Example 1; Page 22; 55pp; English.

XX The present sequence is that of a double-stranded probe for
CC transcription factor Sp-1, which was used in an electrophoretic
CC mobility shift assay in an example from the invention examining
CC prostate cancer-induced expression from the human osteocalcin
CC (hOC) promoter. The cis-acting element OSE-1 was shown to be a
CC weak Sp-1 binding site. An evaluation of the hOC promoter was
CC conducted in which the functional hierarchy of the elements OSE1,
CC OSE2 and AP-1/VDRE (vitamin D response element) was defined in
CC androgen-independent human prostate cancer cell line PC-3. By
CC juxtaposing dimers of these 3 elements, a minimal hOC super-promoter
CC (see ABZ58130) was produced, which displayed over 8-fold higher
CC activity than the native hOC promoter in a tissue-specific manner
CC in PC-3 cells. In one embodiment of the invention, the hOC

CC super-promoter is operably linked to a nucleic acid encoding a
CC heterologous protein, ribozyme, dominant-negative or antisense
CC RNA and used to deliver therapeutic genes to localised or
CC disseminated tumours. hOC promoters can also be used to deliver
CC therapeutic genes to fractured bones for bone repair. hOC promoter
CC activation by extracellular matrices and soluble factors secreted
CC by prostate cancer and bone cells in useful as a marker for the
CC diagnosis and prognosis of prostate cancer.

XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

SQ Query Match 100.0%; Score 22; DB 25; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAGC 22
DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 14

AAF61841
ID AAF61841 standard; DNA; 49 BP.

XX AC AAF61841;
XX XX

DT 26-JUL-2001 (first entry)

DE SPI-specific DNA probe Zi-3.

XX Probe; detection; electron donor; electron acceptor; competitive effect;
KW equilibrium constants determination; allele-specific interaction;
KW gene regulation; antigen identification; allosteric inhibitor; ss.

OS Unidentified.

XX WO200131057-A2.

PN 03-MAY-2001.

PD 17-OCT-2000; 2000WO-EP10209.

PF 22-OCT-1999; 99DE-1050969.

PR (AVET) AVENTIS RES & TECHNOLOGIES GMBH & CO KG.

PA Muth J, Windhab N;
PI WPI; 2001-335702/35.

DR New nucleic acid probe bound to electrically conductive surface, used
PT e.g. to detect antigens or allosteric inhibitors, is attached to an
PT electron donor or acceptor -

PS Example 6; Page 56; 57pp; German.

XX This invention describes a novel nucleic acid probe (I) which comprises
CC a sequence, at least partly double-stranded (ds), bonded to a conductive
CC surface and having attached to it at least one electron donor (ED) or at
CC least one electron acceptor (EA). (I) are used to detect any type of
CC interaction, direct or indirect, that involves (I), also for
CC determination of equilibrium constants, competitive effects and rates of
CC reaction. Some typical applications are identification of antigens and
CC allosteric inhibitors; studying gene regulation (expression or
CC transcription), including allele-specific interactions between
CC regulatory factors and DNA sequences, also separation of interacting
CC factors. (I) are electrically 'readable' and provide rapid and simple
CC detection of any type of (in)direct interaction. Very small changes in
CC conduction can be detected. (I) have good compatibility with a variety
CC of reaction media, e.g. crude extracts can be analyzed. This sequence
CC represents a SPI-specific DNA probe, Zi-3 which is used to illustrate the
CC method of the invention.

SQ Sequence 49 BP; 6 A; 16 C; 16 G; 11 T; 0 other;

Query Match 100.0%; Score 22; DB 22; Length 49;
 Best Local Similarity 100.0%; Pred. No. 3.1;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 ATTCGATCGGGCGGGCGGAGC 22
 ||||||||||||||||||
 DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 15

AAF61841/c

ID AAF61841 standard; DNA; 49 BP.

XX AC AAF61841;

XX DT 26-JUL-2001 (first entry)

XX DE SPI-specific DNA probe Zi-3.

XX KW Probe; detection; electron donor; electron acceptor; competitive effect;

XX KW equilibrium constants determination; allele-specific interaction;

XX KW gene regulation; antigen identification; allosteric inhibitor; ss.

XX OS Unidentified.

XX WO200131057-A2.

XX PD 03-MAY-2001.

XX PF 17-OCT-2000; 2000WO-EPI0209.

XX PR 22-OCT-1999; 99DE-1050969.

XX PA (AVET) AVENTIS RES & TECHNOLOGIES GMBH & CO KG.

XX PI Muth J, Windhab N;

XX DR WPI; 2001-335702/35.

XX PT New nucleic acid probe bound to electrically conductive surface, used

XX PT e.g. to detect antigens or allosteric inhibitors, is attached to an

XX PT electron donor or acceptor

XX PS Example 6; Page 56; 57pp; German.

XX CC This invention describes a novel nucleic acid probe (I) which comprises

XX CC a sequence, at least partly double-stranded (ds), bonded to a conductive

XX CC surface and having attached to it at least one electron donor (ED) or at

XX CC least one electron acceptor (EA). (I) are used to detect any type of

XX CC interaction, direct or indirect, that involves (I), also for

XX CC determination of equilibrium constants, competitive effects and rates of

XX CC reaction. Some typical applications are identification of antigens and

XX CC allosteric inhibitors; studying gene regulation (expression or

XX CC transcription), including allele-specific interactions between

XX CC regulatory factors and DNA sequences, also separation of interacting

XX CC factors. (I) are electrically 'readable' and provide rapid and simple

XX CC detection of any type of (indirect) interaction. Very small changes in

XX CC conduction can be detected. (I) have good compatibility with a variety

XX CC of reaction media, e.g. crude extracts can be analyzed. This sequence

XX CC represents a SPI-specific DNA probe, Zi-3 which is used to illustrate the

XX CC method of the invention.

XX SQ Sequence 49 BP; 6 A; 16 C; 16 G; 11 T; 0 other;

Query Match 100.0%; Score 22; DB 22; Length 49;

Best Local Similarity 100.0%; Pred. No. 3.1;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATTCGATCGGGCGGGCGGAGC 22

|||||

DB 49 ATTCGATCGGGCGGGCGGAGC 28

RESULT 16

AAF61843

ID AAF61843 standard; DNA; 49 BP.

XX AC AAF61843;

XX DT 26-JUL-2001 (first entry)

XX DE SPI-specific DNA probe Zi-5.

XX KW Probe; detection; electron donor; electron acceptor; competitive effect;

XX KW equilibrium constants determination; allele-specific interaction;

XX KW gene regulation; antigen identification; allosteric inhibitor; ss.

XX OS Unidentified.

XX WO200131057-A2.

XX PD 03-MAY-2001.

XX PF 17-OCT-2000; 2000WO-EPI0209.

XX PR 22-OCT-1999; 99DE-1050969.

XX PA (AVET) AVENTIS RES & TECHNOLOGIES GMBH & CO KG.

XX PI Muth J, Windhab N;

XX DR WPI; 2001-335702/35.

XX PT New nucleic acid probe bound to electrically conductive surface, used

XX PT e.g. to detect antigens or allosteric inhibitors, is attached to an

XX PT electron donor or acceptor

XX PS Example 6; Page 56; 57pp; German.

XX CC This invention describes a novel nucleic acid probe (I) which comprises

XX CC a sequence, at least partly double-stranded (ds), bonded to a conductive

XX CC surface and having attached to it at least one electron donor (ED) or at

XX CC least one electron acceptor (EA). (I) are used to detect any type of

XX CC interaction, direct or indirect, that involves (I), also for

XX CC determination of equilibrium constants, competitive effects and rates of

XX CC reaction. Some typical applications are identification of antigens and

XX CC allosteric inhibitors; studying gene regulation (expression or

XX CC transcription), including allele-specific interactions between

XX CC regulatory factors and DNA sequences, also separation of interacting

XX CC factors. (I) are electrically 'readable' and provide rapid and simple

XX CC detection of any type of (indirect) interaction. Very small changes in

XX CC conduction can be detected. (I) have good compatibility with a variety

XX CC of reaction media, e.g. crude extracts can be analyzed. This sequence

XX CC represents a SPI-specific DNA probe, Zi-5 which is used to illustrate the

XX CC method of the invention.

XX SQ Sequence 49 BP; 6 A; 16 C; 16 G; 11 T; 0 other;

Query Match 100.0%; Score 22; DB 22; Length 49;

Best Local Similarity 100.0%; Pred. No. 3.1;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATTCGATCGGGCGGGCGGAGC 22

|||||

DB 1 ATTCGATCGGGCGGGCGGAGC 22

RESULT 17

AAF61843/c

ID AAF61843 standard; DNA; 49 BP.

XX AC AAF61843;

XX DT 26-JUL-2001 (first entry)

XX DE SPI-specific DNA probe Zi-5.

XX KW Probe; detection; electron donor; electron acceptor; competitive effect;

XX KW equilibrium constants determination; allele-specific interaction;

XX KW gene regulation; antigen identification; allosteric inhibitor; ss.

XX OS Unidentified.

XX PN WO200131057-A2.

XX PD 03-MAY-2001.

XX PF 17-OCT-2000; 2000WO-BF10209.

XX PR 22-OCT-1999; 99DE-1050969.

XX PS (AVET) AVENTIS RES & TECHNOLOGIES GMBH & CO KG.

XX PI Muth J, Windhab N;

XX DR WPI; 2001-335702/35.

XX PT New nucleic acid probe bound to electrically conductive surface, used

XX PT e.g. to detect antigens or allosteric inhibitors, is attached to an

XX PT electron donor or acceptor -

XX PS Example 6; Page 56; 57pp; German.

XX CC This invention describes a novel nucleic acid probe (I) which comprises

XX CC a sequence, at least partly double-stranded (ds), bonded to a conductive

XX CC surface and having attached to it at least one electron donor (ED) or at

XX CC least one electron acceptor (EA). (I) are used to detect any type of

XX CC interaction, direct or indirect, that involves (I), also for

XX CC determination of equilibrium constants, competitive effects and rates of

XX CC reaction. Some typical applications are identification of antigens and

XX CC allosteric inhibitors; studying gene regulation (expression or

XX CC transcription), including allele-specific interactions between

XX CC regulatory factors and DNA sequences, also separation of interacting

XX CC factors. (I) are electrically 'readable' and provide rapid and simple

XX CC detection of any type of (in)direct interaction. Very small changes in

XX CC conduction can be detected. (I) have good compatibility with a variety

XX CC of reaction media, e.g. crude extracts can be analyzed. This sequence

XX CC represents a SPI-specific DNA probe, Zi-5 which is used to illustrate the

XX CC method of the invention.

XX SQ Sequence 49 BP; 6 A; 16 C; 16 G; 11 T; 0 other;

Query Match 100.0%; Score 22; DB 22; Length 49;

Best Local Similarity 100.0%; Pred. No. 3.1; Mismatches 0; Indels 0; Gaps 0;

Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGAGC 22

DB 49 ATTCGATCGGGCGGGCGAGC 28

RESULT 18

AAF99620

ID AAF99620 standard; DNA; 21 BP.

AC AAF99620;

XX 12-JUN-2001 (first entry)

XX Immunostimulatory nucleic acid #736.

XX Vaccine; cytostatic; virucidal; bactericidal; fungicidal; anti-parasitic;

XX immunostimulatory; tumour; viral infection; bacterial infection;

XX fungal infection; parasitic infection; cancer; asthma;

XX infectious disease; allergy; immune deficiency; phosphorothioate; ss.

XX OS Synthetic.

XX PN WO200122972-A2.

XX PD 05-APR-2001.

XX PF 25-SEP-2000; 2000WO-US26383.

XX PR 25-SEP-1999; 99US-0156113.

XX PR 27-SEP-1999; 99US-0156135.

XX PR 23-AUG-2000; 2000US-0227436.

XX XX (IOWA) UNIV IOWA RES FOUND.

XX PA (COLE-) COLEY PHARM GMBH.

XX PI Krieg AM, Schetter C, Vollmer J;

XX XX WPI; 2001-273485/28.

XX DR Vaccinating against tumors, infectious diseases, allergies and asthma

XX PT using immunostimulatory Py-rich and TG nucleic acids -

XX XX Claim 101; Page 54; 338pp; English.

XX CC The present invention relates to a method for stimulating an immune

XX CC response. The method comprises administering an immunostimulatory nucleic

XX CC acid to a non-rodent subject in sufficient quantity to stimulate an

XX CC immune response. The present sequence is one such immunostimulatory

XX CC nucleic acid. The immunostimulatory nucleic acids can be pyrimidine rich

XX CC (py-rich) or thymidine (T) rich. The method is used to vaccinate subjects

XX CC against tumour antigens, viral antigens (e.g. herpesviridae, retroviridae

XX CC and/or orthomyxoviridae), bacterial antigens (e.g. toxoplasma,

XX CC haemophilus, campylobacter, clostridium, Escherichia coli and/or

XX CC staphylococcus), fungal antigens and/or parasitic antigens. The method is

XX CC also useful for preventing cancer, asthma, infectious disease, allergy or

XX CC immune deficiency. The present sequence can also be used to redirect a

XX CC Th2 to a Th1 immune response and to activate immune cells.

XX CC Note: the present sequence may have a phosphorothioate backbone.

XX SQ Sequence 21 BP; 3 A; 4 C; 11 G; 3 T; 0 other;

Query Match 95.5%; Score 21; DB 22; Length 21;

Best Local Similarity 100.0%; Pred. No. 8.6; Mismatches 0; Indels 0; Gaps 0;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGAG 21

DB 1 ATTCGATCGGGCGGGCGAG 21

RESULT 19

AAF99621/c

ID AAF99621 standard; DNA; 21 BP.

XX AAF99621;

XX 12-JUN-2001 (first entry)

XX Immunostimulatory nucleic acid #737.

XX Vaccine; cytostatic; virucidal; bactericidal; fungicidal; anti-parasitic;

XX immunostimulatory; tumour; viral infection; bacterial infection;

XX fungal infection; parasitic infection; cancer; asthma;

XX infectious disease; allergy; immune deficiency; phosphorothioate; ss.

XX OS Synthetic.

XX PN WO200122972-A2.

XX PD 05-APR-2001.

XX PF 25-SEP-2000; 2000WO-US26383.

XX PR 25-SEP-1999; 99US-0156113.

PR 27-SEP-1999; 99US-0156135.
 PR 23-AUG-2000; 2000US-0227436.
 XX (IOWA) UNIV IOWA RES FOUND.
 XX (COLE-) COLEY PHARM GMBH.
 XX Krieg AM, Schetter C, Vollmer J;
 XX WPI; 2001-273485/28.
 DR WPI; 2001-273485/28.
 XX Vaccinating against tumors, infectious diseases, allergies and asthma
 XX using immunostimulatory Py-rich and TG nucleic acids -
 XX Claim 101; Page 54; 338pp; English.
 XX The present invention relates to a method for stimulating an immune
 CC response. The method comprises administering an immunostimulatory nucleic
 CC acid to a non-rodent subject in sufficient quantity to stimulate an
 CC immune response. The present sequence is one such immunostimulatory
 CC nucleic acid. The immunostimulatory nucleic acids can be pyrimidine rich
 CC (py-rich) or thymidine (T) rich. The method is used to vaccinate subjects
 CC against tumour antigens, viral antigens (e.g. herpesviridae, retroviridae
 CC and/or orthomyxoviridae), bacterial antigens (e.g. toxoplasma,
 CC haemophilus, campylobacter, clostridium, Escherichia coli and/or
 CC staphylococcus), fungal antigens and/or parasitic antigens. The method is
 CC also useful for preventing cancer, asthma, infectious disease, allergy or
 CC immune deficiency. The present sequence can also be used to redirect a
 CC Th2 to a Th1 immune response and to activate immune cells.
 CC Note: the present sequence may have a phosphorothioate backbone.
 XX Sequence 21 BP; 3 A; 11 C; 4 G; 3 T; 0 other;
 SQ Query Match 95.5%; Score 21; DB 22; Length 21;
 Best Local Similarity 100.0%; Pred. No. 8.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATTCGATCGGGCGGGCGGAG 21
 Db 21 ATTCGATCGGGCGGGCGGAG 1
 RESULT 20
 ABS78341
 ID ABS78341 standard; DNA; 21 BP.
 XX ABS78341;
 AC ABS78341;
 DT 13-DEC-2002 (first entry)
 XX Angiogenesis inhibitory oligonucleotide #825.
 XX Angiogenesis inhibitor; ss; angiogenesis; solid tumour growth;
 KW tumour metastasis; precancerous lesion; rheumatoid arthritis;
 KW psoriasis; diabetic retinopathy; retinopathy of prematurity;
 KW macular degeneration; corneal graft rejection; neovascular glaucoma;
 KW retrolental fibroplasia; rubeosis; Osler-Webber Syndrome;
 KW myocardial angiogenesis; plaque neovascularisation; telangiectasia;
 KW haemophilic joint; angiofibroma; wound granulation;
 KW intestinal adhesion; atherosclerosis; scleroderma; hypertrophic scar.
 XX Synthetic.
 XX WO200253141-A2.
 XX 11-JUL-2002.
 XX 14-DEC-2001; 2001WO-US48458.
 XX 14-DEC-2000; 2000US-255534P.
 XX (COLE-) COLEY PHARM GROUP INC.
 XX Bratzler RL;
 XX WPI; 2002-566690/60.
 XX Inhibiting angiogenesis in a subject, involves administering at least
 XX one antiangiogenic nucleic acid molecule to the subject -
 XX Claim 2; Page 34; 276pp; English.

XX WPI; 2002-566690/60.
 DR Inhibiting angiogenesis in a subject, involves administering at least
 XX one antiangiogenic nucleic acid molecule to the subject -
 XX Claim 2; Page 34; 276pp; English.
 XX The invention relates to inhibiting angiogenesis in a subject, comprising
 CC administering at least one antiangiogenic nucleic acid molecule.
 CC Also included is a kit comprising a first container housing the
 CC antiangiogenic nucleic acids, and instructions for administering them to
 CC a subject having a condition characterised by unwanted angiogenesis.
 CC The method is useful for inhibiting angiogenesis associated with solid
 CC tumour growth, tumour metastasis, precancerous lesion, rheumatoid
 CC arthritis, psoriasis, diabetic retinopathy, retinopathy of prematurity,
 CC macular degeneration, corneal graft rejection, neovascular glaucoma,
 CC retrolental fibroplasia, rubeosis, Osler-Webber Syndrome, myocardial
 CC angiogenesis, plaque neovascularisation, telangiectasia, haemophilic
 CC joints, angiofibroma, wound granulation, intestinal adhesions,
 CC atherosclerosis, scleroderma and hypertrophic scars. The present
 CC sequence is an antiangiogenic nucleic acid of the invention.
 XX Sequence 21 BP; 3 A; 4 C; 11 G; 3 T; 0 other;
 SQ Query Match 95.5%; Score 21; DB 24; Length 21;
 Best Local Similarity 100.0%; Pred. No. 8.6;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATTCGATCGGGCGGGCGGAG 21
 Db 1 ATTCGATCGGGCGGGCGGAG 21
 RESULT 21
 ABS78342/C
 ID ABS78342 standard; DNA; 21 BP.
 XX ABS78342;
 AC ABS78342;
 XX 13-DEC-2002 (first entry)
 XX Angiogenesis inhibitory oligonucleotide #826.
 XX Angiogenesis inhibitor; ss; angiogenesis; solid tumour growth;
 KW tumour metastasis; precancerous lesion; rheumatoid arthritis;
 KW psoriasis; diabetic retinopathy; retinopathy of prematurity;
 KW macular degeneration; corneal graft rejection; neovascular glaucoma;
 KW retrolental fibroplasia; rubeosis; Osler-Webber Syndrome;
 KW myocardial angiogenesis; plaque neovascularisation; telangiectasia;
 KW haemophilic joint; angiofibroma; wound granulation;
 KW intestinal adhesion; atherosclerosis; scleroderma; hypertrophic scar.
 XX Synthetic.
 XX WO200253141-A2.
 XX 11-JUL-2002.
 XX 14-DEC-2001; 2001WO-US48458.
 XX 14-DEC-2000; 2000US-255534P.
 XX (COLE-) COLEY PHARM GROUP INC.
 XX Bratzler RL;
 XX WPI; 2002-566690/60.
 XX Inhibiting angiogenesis in a subject, involves administering at least
 XX one antiangiogenic nucleic acid molecule to the subject -
 XX Claim 2; Page 34; 276pp; English.

XX The invention relates to inhibiting angiogenesis in a subject, comprising
CC administering at least one antiangiogenic nucleic acid molecule.
CC Also included is a kit comprising a first container housing the
CC antiangiogenic nucleic acids, and instructions for administering them to
CC a subject having a condition characterised by unwanted angiogenesis.
CC The method is useful for inhibiting angiogenesis associated with solid
CC tumour growth, tumour metastasis, precancerous lesion, rheumatoid
CC arthritis, psoriasis, diabetic retinopathy, retinopathy of prematurity,
CC macular degeneration, corneal graft rejection, neovascular glaucoma,
CC retrolental fibroplasia, rubecosis, Osler-Weber Syndrome, myocardial
CC angiogenesis, plaque neovascularisation, telangiectasia, haemophilic
CC joints, angiodioma, wound granulation, intestinal adhesions,
CC atherosclerosis, scleroderma and hypertrophic scars. The present
CC sequence is an antiangiogenic nucleic acid of the invention.
XX
SQ Sequence 21 BP; 3 A; 11 C; 4 G; 3 T; 0 other;
Query Match 95.5%; Score 21; DB 24; Length 21;
Best Local Similarity 100.0%; Pred. No. 8.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 ATTCCGATCGGGCGGGCGGCGAG 21
DB 21 ATTCCGATCGGGCGGGCGGCGAG 1
RESULT 22
ABL38746
ID ABL38746 standard; DNA; 21 BP.
XX
AC ABL38746;
XX
DT 16-APR-2002 (first entry)
XX
DE Immunostimulatory nucleic acid SEQ ID NO: 115.
XX
KW Antibody-induced cell lysis; cancer; immunostimulatory; CD20;
XX angiogenesis; metastasis; cytostatic; ss.
XX Synthetic.
XX
XX WO200197843-A2.
XX
XX 27-DEC-2001.
XX
XX 22-JUN-2001; 2001WO-US20154.
XX
XX 22-JUN-2000; 2000US-213346P.
XX
XX (IOWA) UNIV IOWA RES FOUND.
XX
XX Weiner G, Hartmann G;
XX
XX WPI; 2002-154611/20.
XX
XX Treating or preventing cancer, such as basal cell carcinoma, comprises
PT administering immunostimulatory nucleic acids that induce expression of
PT cell surface antigens and antibodies to a subject having or at risk of
PT developing cancer -
XX
XX Disclosure; Page 124; 312pp; English.
XX
XX The present invention relates to methods for treating or preventing
CC cancer, involving administering to a subject having or at risk of
CC developing cancer immunostimulatory nucleic acids that induce expression
CC of cell surface antigens and antibodies. The methods are useful for
CC treating or preventing cancer such as basal cell carcinoma, bladder
CC cancer, bone cancer, brain and central nervous system (CNS) cancer,
CC breast cancer, cervical cancer, colon and rectum cancer, connective
CC tissue cancer, oesophageal cancer, eye cancer, kidney cancer, larynx
CC cancer, leukaemia, liver cancer, lung cancer, Hodgkin's lymphoma,
CC non-Hodgkin's lymphoma, melanoma, myeloma, oral cavity cancer, ovarian
CC cancer, pancreatic cancer, prostate cancer, rhabdomyosarcoma, skin
CC cancer, stomach cancer, testicular cancer, and uterine cancer. The
CC present sequence is an immunostimulatory oligonucleotide described in
CC the exemplification of the invention.
XX
SQ Sequence 21 BP; 3 A; 11 C; 4 G; 3 T; 0 other;
Query Match 95.5%; Score 21; DB 24; Length 21;
Best Local Similarity 100.0%; Pred. No. 8.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CC cancer, pancreatic cancer, prostate cancer, rhabdomyosarcoma, skin
CC cancer, stomach cancer, testicular cancer, and uterine cancer. The
CC present sequence is an immunostimulatory oligonucleotide described in
CC the exemplification of the invention.
XX
SQ Sequence 21 BP; 3 A; 4 C; 11 G; 3 T; 0 other;
Query Match 95.5%; Score 21; DB 24; Length 21;
Best Local Similarity 100.0%; Pred. No. 8.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 ATTCCGATCGGGCGGGCGGCGAG 21
DB 1 ATTCCGATCGGGCGGGCGGCGAG 21
RESULT 23
ABL38839/c
ID ABL38839 standard; DNA; 21 BP.
XX
AC ABL38839;
XX
DT 16-APR-2002 (first entry)
XX
DE Immunostimulatory nucleic acid SEQ ID NO: 229.
XX
KW Antibody-induced cell lysis; cancer; immunostimulatory; CD20;
XX angiogenesis; metastasis; cytostatic; ss.
XX Synthetic.
XX
XX WO200197843-A2.
XX
XX 27-DEC-2001.
XX
XX 22-JUN-2001; 2001WO-US20154.
XX
XX 22-JUN-2000; 2000US-213346P.
XX
XX (IOWA) UNIV IOWA RES FOUND.
XX
XX Weiner G, Hartmann G;
XX
XX WPI; 2002-154611/20.
XX
XX Treating or preventing cancer, such as basal cell carcinoma, comprises
PT administering immunostimulatory nucleic acids that induce expression of
PT cell surface antigens and antibodies to a subject having or at risk of
PT developing cancer -
XX
XX Disclosure; Page 153; 312pp; English.
XX
XX The present invention relates to methods for treating or preventing
CC cancer, involving administering to a subject having or at risk of
CC developing cancer immunostimulatory nucleic acids that induce expression
CC of cell surface antigens and antibodies. The methods are useful for
CC treating or preventing cancer such as basal cell carcinoma, bladder
CC cancer, bone cancer, brain and central nervous system (CNS) cancer,
CC breast cancer, cervical cancer, colon and rectum cancer, connective
CC tissue cancer, oesophageal cancer, eye cancer, kidney cancer, larynx
CC cancer, leukaemia, liver cancer, lung cancer, Hodgkin's lymphoma,
CC non-Hodgkin's lymphoma, melanoma, myeloma, oral cavity cancer, ovarian
CC cancer, pancreatic cancer, prostate cancer, rhabdomyosarcoma, skin
CC cancer, stomach cancer, testicular cancer, and uterine cancer. The
CC present sequence is an immunostimulatory oligonucleotide described in
CC the exemplification of the invention.
XX
SQ Sequence 21 BP; 3 A; 11 C; 4 G; 3 T; 0 other;
Query Match 95.5%; Score 21; DB 24; Length 21;
Best Local Similarity 100.0%; Pred. No. 8.6;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 24
ABT17254
ID ABT17254 standard; DNA; 21 BP.
XX AC ABT17254;
XX DT 10-APR-2003 (first entry)
XX DE Transcription factor probe - SEQ ID No 81.
XX KW Probe; ss; transcription factor-protein complex; transcription factor;
XX KW drug screening; drug identification; array hybridisation.
XX OS Unidentified.
XX PN WO2002101351-A2.
XX PD 19-DEC-2002.
XX PF 30-MAY-2002; 2002WO-US17408.
XX PR 08-JUN-2001; 2001US-0877243.
XX PR 08-JUN-2001; 2001US-0877403.
XX PR 08-JUN-2001; 2001US-0877705.
XX PR 08-JUN-2001; 2001US-0877738.
XX PR 05-SEP-2001; 2001US-0947274.
XX PA (PANO-) PANOMICS INC.
XX PI Li X;
XX DR WPI; 2003-148829/14.
XX PT Identifying transcription factor-protein complexes, by isolating
XX PT transcription factor complexes from sample based on a specific type of
XX PT factor, and identifying different proteins present in isolated
XX PT complexes -
XX PS Disclosure; Fig 6; 167pp; English.
XX CC The invention comprises a method for identifying complexes between a
XX CC transcription factor and another protein. The invention also comprises a
XX CC method for isolating DNA probes which bind to activated transcription
XX CC factors. The methods of the invention are useful for identifying
XX CC transcription factor-protein interactions. The methods of the invention
XX CC are also useful for facilitating the screening and identification of new
XX CC drugs, characterising their mechanism of action and screening for adverse
XX CC side effects based on drug's impact expression. The present DNA sequence
XX CC represents a probe used in the method of the invention.
XX SQ Sequence 21 BP; 3 A; 4 C; 11 G; 3 T; 0 other;
XX Query Match 95.5%; Score 21; DB 25; Length 21;
XX Best Local Similarity 100.0%; Pred. No. 8.6;
XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 1 ATTCGATCGGGCGGGCGGAG 1

RESULT 25
ABT17255/c
ID ABT17255 standard; DNA; 21 BP.
XX AC ABT17255;
XX DT 10-APR-2003 (first entry)

XX DE Transcription factor probe - SEQ ID No 82.
XX KW Probe; ss; transcription factor-protein complex; transcription factor;
XX KW drug screening; drug identification; array hybridisation.
XX OS Unidentified.
XX PN WO2002101351-A2.
XX PD 19-DEC-2002.
XX PF 30-MAY-2002; 2002WO-US17408.
XX PR 08-JUN-2001; 2001US-0877243.
XX PR 08-JUN-2001; 2001US-0877403.
XX PR 08-JUN-2001; 2001US-0877705.
XX PR 08-JUN-2001; 2001US-0877738.
XX PR 05-SEP-2001; 2001US-0947274.
XX PA (PANO-) PANOMICS INC.
XX PI Li X;
XX DR WPI; 2003-148829/14.
XX PT Identifying transcription factor-protein complexes, by isolating
XX PT transcription factor complexes from sample based on a specific type of
XX PT factor, and identifying different proteins present in isolated
XX PT complexes -
XX PS Disclosure; Fig 6; 167pp; English.
XX CC The invention comprises a method for identifying complexes between a
XX CC transcription factor and another protein. The invention also comprises a
XX CC method for isolating DNA probes which bind to activated transcription
XX CC factors. The methods of the invention are useful for identifying
XX CC transcription factor-protein interactions. The methods of the invention
XX CC are also useful for facilitating the screening and identification of new
XX CC drugs, characterising their mechanism of action and screening for adverse
XX CC side effects based on drug's impact expression. The present DNA sequence
XX CC represents a probe used in the method of the invention.
XX SQ Sequence 21 BP; 3 A; 11 C; 4 G; 3 T; 0 other;
XX Query Match 95.5%; Score 21; DB 25; Length 21;
XX Best Local Similarity 100.0%; Pred. No. 8.6;
XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 26
ABT17322/c
ID ABT17322 standard; DNA; 63 BP.
XX AC ABT17322;
XX DT 10-APR-2003 (first entry)
XX DE Transcription factor-related array hybridisation probe - SEQ ID No 149.
XX KW Probe; ss; transcription factor-protein complex; transcription factor;
XX KW drug screening; drug identification; array hybridisation.
XX OS Unidentified.
XX PN WO2002101351-A2.
XX PD 19-DEC-2002.

PT infections and to modulate immune response e.g. tolerance break and
 PT regulation of TH1/TH2 cells

PS Example 5; Page 9; 28pp; English.

XX AAV45993-V46019 are fragments of bacterial polynucleotides which are
 CC used as immune adjuvants for inclusion into vaccines to treat cancer and
 CC for prophylaxis and/or treatment of conditions caused by pathogenic
 CC micro-organisms. The polynucleotide is used for modulation of an immune
 CC response and the modulation is selected for the group break of
 CC tolerance, regulation of TH1/TH2 helper cell responses, switch of Ig
 CC classes, treatment of autoimmune responses and induction of tolerances.
 CC DNA oligomers are used to enhance the reactivity of immune cells to
 CC viral, bacterial and parasitic antigens, to break tolerance in vaccinee
 CC and B cells e.g. against tumour antigens, as adjuvants in vaccination
 CC against tumour-defined antigens and immunostimulatory substances in an
 CC immune response against tumours and to suppress immune reactions of the
 CC innate and acquired immune system. The composition is inexpensive and
 CC stable and does not cause lethal shock, which happens with prior art
 CC bacterial sequences.

XX Sequence 20 BP; 2 A; 5 C; 11 G; 2 T; 0 other;

Query Match 90.9%; Score 20; DB 19; Length 20;

Best Local Similarity 100.0%; Pred. No. 23;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGCGAGC 22

Db 1 TCGATCGGGCGGGCGGCGAGC 20

RESULT 29

AAZ99627

ID AAZ99627 standard; DNA; 20 BP.

XX AC AAZ99627;

DT 12-JUL-2000 (first entry)

DE Nucleotide sequence of G-motif oligonucleotide SP1.

XX G-motif oligonucleotide; vaccine; Toxoplasmosis; viral infection;
 KW antigen presenting cell activation; natural killer cell; septic shock;
 KW cytotoxic T-lymphocyte; inflammation; autoimmune disease;
 KW rheumatoid arthritis; Crohn's disease; sarcoidosis; multiple sclerosis;
 KW Kawasaki syndrome; graft-versus-host disease; transplant rejection;
 KW helper T cell response i-mediated disease; Lyme arthritis;
 KW Streptococcal induced arthritis; chronic inflammatory bowel disease;
 KW psoriasis vulgaris; experimental allergic encephalomyelitis;
 KW insulin-dependent diabetes mellitus; bacterial infection;
 KW parasitic infection; Leishmaniasis; spontaneous abortion; tumour; ss.

XX Synthetic.

XX WO200014217-A2.

XX 16-MAR-2000.

XX 03-SEP-1999; 99WO-EF06502.

XX 03-SEP-1998; 98EP-0116652.

XX (CPGI-) CPG IMMUNOPHARMACEUTICALS GMBH.

XX Wagner H, Lipford GB, Keeg K;

XX WPI; 2000-256970/22.

XX Compositions comprising G-motif oligonucleotides useful for treating
 PT e.g. septic shock, rheumatoid arthritis, diabetes and human
 PT immunodeficiency virus infections -

PS Example 14; Page 32; 75pp; English.

XX The present sequence represents a G-motif oligonucleotide of the
 CC invention. The specification describes compositions comprising G-motif
 CC oligonucleotides. The G-motif oligonucleotides inhibit activation of
 CC antigen presenting cells by inhibiting the uptake of DNA by a cell, by
 CC stimulating natural killer cells, or by co-stimulating cytotoxic
 CC T-lymphocytes. The G-motif oligonucleotides may be used for the
 CC productions of vaccines for treating septic shock, inflammation,
 CC autoimmune diseases (e.g. rheumatoid arthritis, Crohn's disease,
 CC sarcoidosis, multiple sclerosis, Kawasaki syndrome, graft-versus-host
 CC disease and transplant rejection), helper T cell response i-mediated
 CC diseases (e.g. Streptococcal induced arthritis, Lyme arthritis, chronic
 CC inflammatory bowel disease, psoriasis vulgaris, experimental allergic
 CC encephalomyelitis and insulin-dependent diabetes mellitus), bacterial
 CC infections, parasitic infections and immunostimulatory substances in an
 CC viral infections (e.g. Cytomegalovirus and human immunodeficiency virus
 CC (HIV)-infections), spontaneous abortions and tumours. They may also be
 CC used to induce proliferation of bone marrow cells, especially macrophage
 CC precursor cells.

XX Sequence 20 BP; 2 A; 5 C; 11 G; 2 T; 0 other;

Query Match 90.9%; Score 20; DB 21; Length 20;

Best Local Similarity 100.0%; Pred. No. 23;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGCGAGC 22

Db 1 TCGATCGGGCGGGCGGCGAGC 20

RESULT 30

AAZ99229

ID AAZ99229 standard; DNA; 20 BP.

XX AC AAZ99229;

DT 05-SEP-2002 (first entry)

DE Murine Toll-like receptor related Cpg DNA SEQ ID No 104.

XX Murine Toll-like receptor; TLR9; TLR7; TLR8; ISNA; ds.

XX Unidentified.

XX WO200222809-A2.

XX 21-MAR-2002.

XX 17-SEP-2001; 2001WO-US29229.

XX 15-SEP-2000; 2000US-233035P.

XX 23-JAN-2001; 2001US-263657P.

XX 17-MAY-2001; 2001US-291726P.

XX 22-JUN-2001; 2001US-300210P.

XX (COLE-) COLEY PHARM GMBH.

XX Bauer S, Lipford G, Wagner H;

XX WPI; 2002-393964/42.

XX New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,
 PT useful for identifying species specificity of immunostimulatory nucleic
 PT acid and identifying immunostimulatory nucleic acids -

XX Disclosure; Page 77; 195pp; English.

XX The invention relates to isolated murine Toll-like receptors (TLR)9,
 CC TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined
 CC sequences of 1032, 1050 or 1032 amino acids as given in specification, or
 CC their fragments, where TLR9, TLR7 and TLR8 polypeptides or their

CC fragments have an amino acid sequence which is identical to human TLR9,
 CC TLR7 or TLR8 polypeptides or their fragment except for at least one amino
 CC acid of a murine TLR polypeptide. The isolated nucleic acids of the
 CC invention are useful for inhibiting TLR9 signalling activity in a cell.
 CC TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid
 CC molecules which interact with a TLR polypeptide or its fragment. The
 CC TLR7, TLR8 or TLR9 polypeptides are also useful for identifying ISNA. The
 CC TLR7, TLR8 and TLR9 polypeptides are also useful for comparing TLR9
 CC signalling activity of a test compound (that is not a nucleic acid, and
 CC is a polypeptide or a part of a combinatorial library of compounds) with
 CC an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for
 CC identifying species specificity of an ISNA. The isolated nucleic acids of
 CC the invention are useful as probes or primers. This polynucleotide
 CC sequence represents DNA relating to the isolated Toll-like receptors of
 CC the invention.

SQ Sequence 20 BP; 2 A; 5 C; 11 G; 2 T; 0 other;

Query Match 90.9%; Score 20; DB 24; Length 20;
 Best Local Similarity 100.0%; Pred. No. 23;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 TCGATCGGGCGGGCGGAGC 22
 DB 1 TCGATCGGGCGGGCGGAGC 20

RESULT 32

ABA92272
 ID ABA92272 standard; DNA; 22 BP.

XX
 AC ABA92272;

DT 10-JUN-2002 (first entry)

DE Sp-1 mutant oligonucleotide, used in EMSA.

XX SP-1; neuron; antialzheimers; antiparkinsonian; antisclerotic;
 KW neuroprotective; nootropic; anticonvulsant; vascular; hypotensive;
 KW cerebroprotective; virucide; anti-HIV; diagnosis; therapy;
 KW electrophoretic mobility shift assay; EMSA; ds.

XX Homo sapiens.
 OS Synthetic.

XX WO200215912-A1.

PN 28-FEB-2002.

PD 24-AUG-2001; 2001WO-US26527.

PF 25-AUG-2000; 2000US-228201P.

PR 26-OCT-2000; 2000US-243295P.

XX (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.

PA Ratan RR, Chatterjee S;

PI WPI; 2002-242023/29.

XX Diagnosing and treating diseases associated with oxidative stress, DNA
 PT damage or growth factor depletion, e.g. Alzheimer's and Parkinson's, by
 PT administering e.g. mithramycin, chromomycin, daunomycin, olivomycin and
 PT WP631 -

PS Example 1; Page 23; 69pp; English.

XX The present sequence is that of a double-stranded mutated Sp-1
 CC oligonucleotide, which was used in an electrophoretic mobility
 CC shift assay to determine the effect of oxidative stress on Sp-1 DNA
 CC binding, and to determine the effects of candidate compounds on Sp-1
 CC protein levels. Sp-1 DNA binding activity in cortical neurons was
 CC shown to be low, but was dramatically enhanced by oxidative stress.

CC The invention provides methods for detecting and treating diseases
 CC associated with oxidative stress, DNA damage or growth factor
 CC depletion, and identifying agents for the treatment of such
 CC conditions. A compound is deemed to be an inhibitor of oxidative
 CC stress, DNA damage, growth factor depletion or cell death if it
 CC reduces the protein level of an Sp family member or if it decreases
 CC the binding of an Sp family member to DNA. A method for preventing
 CC or treating a disease or disorder of the nervous system, the ageing
 CC process or associated with apoptosis involves administering a
 CC compound that inhibits the induction of an Sp family member or the
 CC binding of an Sp family member to DNA, e.g. mithramycin,
 CC chromomycin, daunomycin, olivomycin or WP631. Diseases and
 CC disorders that can be treated include Alzheimer's disease, and
 CC Creutzfeldt-Jacob disease, kuru, Huntington's disease, aneurysm,
 CC stroke associated with an increase in blood pressure, spinal cord
 CC disease, spinal cord injury, brain injury, multiple system atrophy,
 CC amyotrophic lateral sclerosis, progressive supranuclear palsy,
 CC neurodegeneration associated with the ageing process, mitochondrial
 CC disease, HIV infection, herpes infection and multiple sclerosis
 CC (all claimed).

XX Sequence 22 BP; 3 A; 5 C; 9 G; 5 T; 0 other;

Query Match 85.5%; Score 18.8; DB 24; Length 22;
 Best Local Similarity 90.9%; Pred. No. 71;

Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 ATTCGATCGGGCGGGCGGAGC 22

DB 1 ATTCGATCGGGTTCGGGCGGAGC 22

RESULT 32

AAQ67304

ID AAQ67304 standard; DNA; 22 BP.

XX AAQ67304;

AC 30-MAR-1995 (first entry)

DT Detection probe for DNA binding activity of Sp1 protein.

DE Jun protein; Fos protein; Sp1 protein; DNA binding protein;
 KW probe; detection; calibration; ss.

XX Synthetic.

OS JP06201692-A.

PN 22-JUL-1994.

PD 22-SEP-1993; 93JP-0257482.

PF 22-SEP-1992; 92JP-0278126.

PR (YAWH) NIPPON STEEL CHEM CO.
 PA (YAWA) NIPPON STEEL CORP.

XX WPI; 1994-273729/34.

XX Probe for detection or calibration of DNA binding - and detection
 PT or calibration of the protein by using it in situ

XX Claim 5; Page 5; 6pp; Japanese.

XX The probes given in AAQ67303-04 are used for the detection or
 CC calibration of DNA binding and contain nucleotide sequences binding
 CC specifically with DNA binding protein (Jun or Fos protein, and Sp1
 CC protein respectively). Distribution of DNA binding protein in
 CC tissues or cells can be inspected in situ, by using DNA binding
 CC activity as indicator.

XX Sequence 22 BP; 3 A; 5 C; 11 G; 3 T; 0 other;

Query Match 83.6%; Score 18.4; DB 15; Length 22;
Best Local Similarity 95.0%; Pred. No. 1e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
|||||
Db 3 TCGATCGGGCGGGCGGATC 22
|||||

RESULT 33
AAQ30483
ID AAQ30483 standard; DNA; 45 BP.
XX AC AAQ30483;
XX DT 25-MAR-2003 (updated)
XX DT 19-MAR-1993 (first entry)
XX DE Oligonucleotide contg. Sp1 recognition sequence.
XX KW Control recognition element; decoy; cellular RNA; dumbbell;
XX KW promoter; hormone receptor element; viral; liver; tissue; viral;
XX KW proliferation; linker; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT misc_structure 1..45
XX FT /*tag= a
XX FT /note= "looped self structure"
XX PN WO9218522-A1.
XX PD 29-OCT-1992.
XX PF 17-APR-1992; 92WO-US03205.
XX PR 18-APR-1991; 91US-0687337.
XX PA (SALK) SALK INST BIOLOGICAL STUDIES.
XX PI Chu BC, Orgel L;
XX DR WPI; 1992-382035/46.
XX PT New oligo-nucleotide(s) contg. transcription control recognition
XX PT element - stabilised by covalent bonding of two DNA strands, act
XX PT as decoys for regulatory protein to modulate specific RNA
XX PS Example 3; Page 27; 41pp; English.
XX CC The oligonucleotide contains the double stranded control recognition
XX CC element Sp1 recognition sequence, and was synthesised in this linear
XX CC form for prodn. of dumbbell oligonucleotides. The oligomer forms
XX CC a looped self structure with a nicked gap between the 3'-OH and 5'-OH
XX CC tail ends. Certain bases may optionally have incorporated a
XX CC phosphorothioate diester linkage instead of the normal phosphodiester
XX CC bond. The oligonucleotide acts as a decoy for proteins which
XX CC exert transcription control, and so modulate specific cellular RNAs.
XX CC Typical CREs which may be regulated include promoters, hormone
XX CC receptor elements, viral, cellular, liver or tissue elements, etc, and
XX CC a typical application is inhibition of viral proliferation.
XX CC See also AAQ30472-518.
XX CC (Updated on 25-MAR-2003 to correct PN field.)
XX SQ Sequence 45 BP; 3 A; 21 C; 14 G; 7 T; 0 other;
Query Match 81.8%; Score 18; DB 13; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGCGA 20
|||||

Db 28 TCGATCGGGCGGGCGGCGA 45
|||||

RESULT 34
AAQ30483/c
ID AAQ30483 standard; DNA; 45 BP.
XX AC AAQ30483;
XX DT 25-MAR-2003 (updated)
XX DT 19-MAR-1993 (first entry)
XX DE Oligonucleotide contg. Sp1 recognition sequence.
XX KW Control recognition element; decoy; cellular RNA; dumbbell;
XX KW promoter; hormone receptor element; viral; liver; tissue; viral;
XX KW proliferation; linker; ss.
XX OS Synthetic.
XX FH Key Location/Qualifiers
XX FT misc_structure 1..45
XX FT /*tag= a
XX FT /note= "looped self structure"
XX PN WO9218522-A1.
XX PD 29-OCT-1992.
XX PF 17-APR-1992; 92WO-US03205.
XX PR 18-APR-1991; 91US-0687337.
XX PA (SALK) SALK INST BIOLOGICAL STUDIES.
XX PI Chu BC, Orgel L;
XX DR WPI; 1992-382035/46.
XX PT New oligo-nucleotide(s) contg. transcription control recognition
XX PT element - stabilised by covalent bonding of two DNA strands, act
XX PT as decoys for regulatory protein to modulate specific RNA
XX PS Example 3; Page 27; 41pp; English.
XX CC The oligonucleotide contains the double stranded control recognition
XX CC element Sp1 recognition sequence, and was synthesised in this linear
XX CC form for prodn. of dumbbell oligonucleotides. The oligomer forms
XX CC a looped self structure with a nicked gap between the 3'-OH and 5'-OH
XX CC tail ends. Certain bases may optionally have incorporated a
XX CC phosphorothioate diester linkage instead of the normal phosphodiester
XX CC bond. The oligonucleotide acts as a decoy for proteins which
XX CC exert transcription control, and so modulate specific cellular RNAs.
XX CC Typical CREs which may be regulated include promoters, hormone
XX CC receptor elements, viral, cellular, liver or tissue elements, etc, and
XX CC a typical application is inhibition of viral proliferation.
XX CC See also AAQ30472-518.
XX CC (Updated on 25-MAR-2003 to correct PN field.)
XX SQ Sequence 45 BP; 3 A; 21 C; 14 G; 7 T; 0 other;
Query Match 81.8%; Score 18; DB 13; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGGAGC 22
|||||
Db 24 GATCGGGCGGGCGGAGC 7
|||||

RESULT 35
AAQ30484/c

```
ID ARQ30484 standard; DNA; 47 BP.
XX
AC AAQ30484;
XX
DT 25-MAR-2003 (updated)
DT 19-MAR-1993 (first entry)
XX
DE Oligonucleotide contg. Spt recognition sequence.
XX
KW Control recognition element; decoy; cellular RNA; dumbbell;
KW promoter; hormone receptor element; viral; liver; tissue; viral;
KW proliferation; linker; cyclic; closed circular.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT misc_structure 1..47
FT /*tag= a
FT /note= "closed, circular, self complementary
FT structure"
XX
PN WO9218522-A1.
XX
XX 29-OCT-1992.
XX
PD 17-APR-1992; 92WO-US03205.
XX
PR 18-APR-1991; 91US-0687337.
XX
PA (SALK ) SALK INST BIOLOGICAL STUDIES.
XX
PI Chu BC, Orgel L;
XX
DR WPI; 1992-382035/46.
XX
XX New oligo-nucleotide(s) contg. transcription control recognition
PT element - stabilised by covalent bonding of two DNA strands, act
PT as decoys for regulatory protein to modulate specific RNA
XX
PS Example 3; Page 27; 41pp; English.
XX
CC The oligonucleotide contains the double stranded control recognition
CC element Spt recognition sequence in the form of a dumbbell structure.
CC The oligomer, when synthesised in its linear form forms a looped
CC self structure with a nicked gap between the 3'-OH and 5'-OH tail
CC ends, which may be phosphorylated with T4 polynucleotide kinase and
CC ligated with DNA ligase to give a closed circular self complementary
CC structure. The oligonucleotide acts as a decoy for proteins which
CC exert transcription control, and so modulate specific cellular RNAs.
CC Typical CREs which may be regulated include promoters, hormone
CC receptor elements, viral, cellular, liver or tissue elements, etc, and
CC a typical application is inhibition of viral proliferation.
CC See also AAQ30472-519.
CC (Updated on 25-MAR-2003 to correct PN field.)
XX
SQ Sequence 47 BP; 3 A; 21 C; 16 G; 7 T; 0 other;
Query Match 81.8%; Score 18; DB 13; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 5 GATCGGGGGGGGGCGAGC 22
DB 42 GATCGGGGGGGGGCGAGC 25
RESULT 36
ID ABA00746/c
XX ABA00746 standard; DNA; 183 BP.
AC ABA00746;
XX
DT 18-MAR-2003 (first entry)
```

```
XX Enhl enhancer.
DE
XX
KW Enhancer; Enhl; lentivirus; producer cell; gag; pol; tissue-specific;
KW transfer vector; blood clotting factor; tat; bleeding disorder; env;
KW hepatocyte transcription factor; HNF1; HNF3; HNF4; HNF6; haemophilia;
KW liver-specific; transgene delivery; gene therapy; ESP; ss.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT misc_RNA 11..161
FT /*tag= a
FT /note= "Hepatocyte derived elements"
XX
PN WO200292134-A1.
XX
XX 21-NOV-2002.
XX
PF 14-MAY-2002; 2002WO-US15062.
XX
PR 14-MAY-2001; 2001US-291083P.
XX
XX (CELL-) CELL GENESYS INC.
XX
XX McArthur JG, Talbot DJ, Simmons AD, McGuinness R, Kelly M;
PI Tsui LV, Dull T;
XX
XX WPI; 2003-120615/11.
XX
XX New lentiviral producer cell, useful for optimizing liver-specific
XX transgene delivery for treating bleeding disorders, e.g. haemophilia, or
XX for tissue-specific gene therapy -
Claim 11; Page 39; 61pp; English.
XX
CC This sequence represents a cloned enhancer element (Enhl) which was
CC used in the production of the lentiviral producer cell of the invention.
CC The producer cell comprises:
CC (a) a first nucleotide sequence comprising a gag, a pol, or gag and pol
CC genes;
CC (b) a second nucleotide sequence comprising a heterologous env gene;
CC and
CC (c) a third nucleotide sequence comprising a lentiviral transfer vector
CC that comprises a gene that encodes a blood clotting factor operably
CC linked to an expression control sequence. The producer cell lacks a
CC functional tat gene. This enhancer sequence was created by ligating
CC together five synthetic oligonucleotide binding sites for hepatocyte
CC transcription factors. This enhancer element augmented expression
CC levels up to two orders of magnitude over the enhancer-less vector. Enhl
CC comprises the following elements:
CC HNF1(sense) - HNF3(sense) - HNF4(antisense) - HNF4(antisense) -
CC HNF6(sense) - EBP(antisense) - HNF4(antisense).
CC The lentiviral producer cell and transfer vectors are useful for
CC optimizing liver-specific transgene delivery for treating bleeding
CC disorders, e.g. haemophilia. The vectors are also useful for tissue-
CC specific gene therapy.
XX
SQ Sequence 183 BP; 40 A; 53 C; 54 G; 36 T; 0 other;
Query Match 79.1%; Score 17.4; DB 25; Length 183;
Best Local Similarity 94.7%; Pred. No. 2.3e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY 4 CGATCGGGGGGGGGCGAGC 22
DB 183 CGATCGGGGGGGGGCGAAC 165
RESULT 37
ID ABAQ44940
XX ABAQ44940 standard; DNA; 722 BP.
XX
```

AC ABQ44940;
XX
XX
XX 12-JUL-2002 (first entry)
XX
XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 31531.
XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.
XX Homo sapiens.
XX
XX WO200218632-A2.
XX
XX 07-MAR-2002.
XX
XX 01-SEP-2001; 2001WO-EPI0074.
XX
XX 01-SEP-2000; 2000DE-1043826.
XX
XX 05-SEP-2000; 2000DE-1044543.
XX
XX (EPIG-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guetig D;
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful
XX for diagnosis and prognosis, comprises selective hybridization of
XX amplicons from chemically treated DNA -
XX
XX Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
XX methylation of a particular cytosine in a motif 5'-CpG-3', present in a
XX genomic sample of DNA. The sample is treated chemically to convert
XX cytosine (C) but not methylated C, to uracil, then part of the genomic
XX DNA that contains the target C is amplified to form a labeled amplicon.
XX The amplicon is hybridised to two classes, each with at least one
XX member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
XX and the degree of hybridisation to both classes is determined from the
XX label on the amplicon. From the ratio of labels hybridised to the two
XX classes of oligomers, the degree of methylation is calculated. The method
XX is used: (i) for diagnosis and/or prognosis of side effects of
XX therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
XX of the central nervous, cardiovascular, gastrointestinal and respiratory
XX systems etc., particularly by detecting mutations or single nucleotide
XX polymorphisms (SNP's); and (ii) for differentiation of cell or tissue
XX types and for investigating cell differentiation. The method allows the
XX methylation status of many C residues to be determined simultaneously.
XX ABQ44941-ABQ44942 represent genomic DNA sequences used to illustrate the
XX method for determining the degree of cytosine methylation described in
XX the disclosure of the invention.
XX
XX Sequence 722 BP; 80 A; 93 C; 286 G; 263 T; 0 other;
SQ
Query Match 79.1%; Score 17.4; DB 24; Length 722;
Best Local Similarity 94.7%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 ATTTCGATCGGGCGGGCGG 19
DB 514 ATTTCGATCGGGCGGGCGG 532
RESULT 38
ABQ44941/c
ID ABQ44941 standard; DNA; 722 BP.
XX
XX AC ABQ44941;
XX
XX 12-JUL-2002 (first entry)
XX
XX

XX
XX
XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 31532.
XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.
XX Homo sapiens.
XX
XX WO200218632-A2.
XX
XX 07-MAR-2002.
XX
XX 01-SEP-2001; 2001WO-EPI0074.
XX
XX 01-SEP-2000; 2000DE-1043826.
XX
XX 05-SEP-2000; 2000DE-1044543.
XX
XX (EPIG-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guetig D;
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful
XX for diagnosis and prognosis, comprises selective hybridization of
XX amplicons from chemically treated DNA -
XX
XX Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
XX methylation of a particular cytosine in a motif 5'-CpG-3', present in a
XX genomic sample of DNA. The sample is treated chemically to convert
XX cytosine (C) but not methylated C, to uracil, then part of the genomic
XX DNA that contains the target C is amplified to form a labeled amplicon.
XX The amplicon is hybridised to two classes, each with at least one
XX member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
XX and the degree of hybridisation to both classes is determined from the
XX label on the amplicon. From the ratio of labels hybridised to the two
XX classes of oligomers, the degree of methylation is calculated. The method
XX is used: (i) for diagnosis and/or prognosis of side effects of
XX therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
XX of the central nervous, cardiovascular, gastrointestinal and respiratory
XX systems etc., particularly by detecting mutations or single nucleotide
XX polymorphisms (SNP's); and (ii) for differentiation of cell or tissue
XX types and for investigating cell differentiation. The method allows the
XX methylation status of many C residues to be determined simultaneously.
XX ABQ44940-ABQ44941 represent genomic DNA sequences used to illustrate the
XX method for determining the degree of cytosine methylation described in
XX the disclosure of the invention.
XX
XX Sequence 722 BP; 263 A; 286 C; 93 G; 80 T; 0 other;
SQ
Query Match 79.1%; Score 17.4; DB 24; Length 722;
Best Local Similarity 94.7%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 ATTTCGATCGGGCGGGCGG 19
DB 209 ATTTCGATCGGGCGGGCGG 191
RESULT 39
ABQ43520
ID ABQ43520 standard; DNA; 747 BP.
XX
XX AC ABQ43520;
XX
XX 12-JUL-2002 (first entry)
XX
XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 30111.
XX
XX

Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis; drug; side effect; cancer; central nervous system; cardiovascular; gastrointestinal; respiratory system; single nucleotide polymorphism; SNP; cell differentiation; ds.

OS Homo sapiens.
 XX WO200218632-A2.
 XX
 PD 07-MAR-2002.
 XX
 PF 01-SEP-2001; 2001WO-EP10074.
 XX
 PR 01-SEP-2000; 2000DE-1043826.
 PR 05-SEP-2000; 2000DE-1044543.
 XX
 XX (EPIG-) EPIGENOMICS AG.
 PI Olek A, Piepenbrock C, Berlin K, Guetig D;
 XX WPI; 2002-371829/40.
 XX
 PT Determining the degree of cytosine methylation in genomic DNA, useful
 PT for diagnosis and prognosis, comprises selective hybridization of
 PT amplicons from chemically treated DNA -
 XX
 PS Claim 12; 56pp + Sequence Listing; 56pp; German.
 XX
 CC This invention describes a novel method for determining the degree of
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
 CC genomic sample of DNA. The sample is treated chemically to convert
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic
 CC DNA that contains the target C is amplified to form a labeled amplicon.
 CC The amplicon is hybridised to two classes, each with at least one
 CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
 CC and the degree of hybridisation to both classes is determined from the
 CC label on the amplicon. From the ratio of labels hybridised to the two
 CC classes of oligomers, the degree of methylation is calculated. The method
 CC is used: (i) for diagnosis and/or prognosis of side effects of
 CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
 CC of the central nervous, cardiovascular, gastrointestinal and respiratory
 CC systems etc., particularly by detecting mutations or single nucleotide
 CC polymorphisms (SNPs); and (ii) for differentiation of cell or tissue
 CC types and for investigating cell differentiation. The method allows the
 CC methylation status of many C residues to be determined simultaneously.
 CC ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the
 CC method for determining the degree of cytosine methylation described in
 CC the disclosure of the invention.
 XX
 SQ Sequence 747 BP; 80 A; 98 C; 302 G; 265 T; 2 other;

Query Match 79.1%; Score 17.4; DB 24; Length 747;
 Best Local Similarity 94.7%; Pred. No. 2.1e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCG 19
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 DB 539 ATTCGATCGGGCGGGCG 557

RESULT 40
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 ID ABQ43521 standard; DNA; 747 BP.
 XX AC ABQ43521;
 XX
 XX 12-JUL-2002 (first entry)
 XX
 DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 30112.
 XX
 KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
 KW drug; side effect; cancer; central nervous system; cardiovascular;
 KW gastrointestinal; respiratory system; single nucleotide polymorphism;

SNP; cell differentiation; ds.

OS Homo sapiens.
 XX WO200218632-A2.
 XX
 PD 07-MAR-2002.
 XX
 PF 01-SEP-2001; 2001WO-EP10074.
 XX
 PR 01-SEP-2000; 2000DE-1043826.
 PR 05-SEP-2000; 2000DE-1044543.
 XX
 XX (EPIG-) EPIGENOMICS AG.
 PI Olek A, Piepenbrock C, Berlin K, Guetig D;
 XX WPI; 2002-371829/40.
 XX

Determining the degree of cytosine methylation in genomic DNA, useful
 for diagnosis and prognosis, comprises selective hybridization of
 amplicons from chemically treated DNA -

Claim 12; 56pp + Sequence Listing; 56pp; German.

This invention describes a novel method for determining the degree of
 methylation of a particular cytosine in a motif 5'-CpG-3', present in a
 genomic sample of DNA. The sample is treated chemically to convert
 cytosine (C) but not methylated C, to uracil, then part of the genomic
 DNA that contains the target C is amplified to form a labeled amplicon.
 The amplicon is hybridised to two classes, each with at least one
 member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
 and the degree of hybridisation to both classes is determined from the
 label on the amplicon. From the ratio of labels hybridised to the two
 classes of oligomers, the degree of methylation is calculated. The method
 is used: (i) for diagnosis and/or prognosis of side effects of
 therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
 of the central nervous, cardiovascular, gastrointestinal and respiratory
 systems etc., particularly by detecting mutations or single nucleotide
 polymorphisms (SNPs); and (ii) for differentiation of cell or tissue
 types and for investigating cell differentiation. The method allows the
 methylation status of many C residues to be determined simultaneously.
 ABQ13410-ABQ54121 represent genomic DNA sequences used to illustrate the
 method for determining the degree of cytosine methylation described in
 the disclosure of the invention.

Sequence 747 BP; 265 A; 302 C; 98 G; 80 T; 2 other;

Query Match 79.1%; Score 17.4; DB 24; Length 747;
 Best Local Similarity 94.7%; Pred. No. 2.1e+02;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCG 19
 |||||
 DB 209 ATTCGATCGGGCGGGCG 191

Search completed: February 18, 2004, 16:45:33
 Job time : 173 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 14:54:35 ; Search time 32 seconds
(without alignments)
303.451 Million cell updates/sec

Title: US-10-026-341a-2

Perfect score: 22

Sequence: 1 attcgatcggggcgagcgcgagc 22

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 569978 seqs, 220691566 residues

Total number of hits satisfying chosen parameters: 1139956

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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2: /cgn2_6/ptodata/1/ina/5B_COMB.seq.*
3: /cgn2_6/ptodata/1/ina/6A_COMB.seq.*
4: /cgn2_6/ptodata/1/ina/6B_COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PCTUS_COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----|--------------------|
| 1 | 22 | 100.0 | 22 | 2 | US-08-327-832-12 |
| 2 | 22 | 100.0 | 22 | 2 | US-08-828-584-12 |
| 3 | 22 | 100.0 | 22 | 5 | PCT-US94-05659-16 |
| 4 | 19 | 86.4 | 46 | 1 | US-08-122-433-12 |
| 5 | 19 | 86.4 | 46 | 1 | US-08-122-433-13 |
| 6 | 18 | 81.8 | 46 | 1 | US-08-122-433-12 |
| 7 | 18 | 81.8 | 46 | 1 | US-08-122-433-13 |
| 8 | 16.8 | 76.4 | 2875 | 3 | US-08-458-434A-4 |
| 9 | 16.8 | 76.4 | 15144 | 3 | US-08-458-434A-6 |
| 10 | 16.4 | 74.5 | 379 | 1 | US-09-591-383-5 |
| 11 | 16.4 | 74.5 | 379 | 1 | US-08-145-617-5 |
| 12 | 16.2 | 73.6 | 3997 | 4 | US-09-345-236B-145 |
| 13 | 16.2 | 73.6 | 3997 | 4 | US-09-345-236B-146 |
| 14 | 16.2 | 73.6 | 9208 | 4 | US-09-068-506-1 |
| 15 | 15.8 | 71.8 | 31 | 1 | US-08-153-563-4 |
| 16 | 15.8 | 71.8 | 31 | 2 | US-09-038-227-9 |
| 17 | 15.8 | 71.8 | 31 | 2 | US-08-460-507-4 |
| 18 | 15.8 | 71.8 | 944 | 2 | US-08-786-606-4 |
| 19 | 15.8 | 71.8 | 1112 | 2 | US-08-333-750C-97 |
| 20 | 15.8 | 71.8 | 1112 | 3 | US-09-234-613-97 |
| 21 | 15.8 | 71.8 | 1114 | 4 | US-09-690-454-39 |
| 22 | 15.8 | 71.8 | 1919 | 1 | US-07-991-587A-1 |
| 23 | 15.8 | 71.8 | 1919 | 1 | US-08-309-985-1 |
| 24 | 15.8 | 71.8 | 3314 | 1 | US-07-973-324B-5 |
| 25 | 15.8 | 71.8 | 3314 | 1 | US-08-343-380-5 |
| 26 | 15.8 | 71.8 | 3314 | 3 | US-09-072-435-5 |
| 27 | 15.8 | 71.8 | 3314 | 3 | US-09-072-917A-5 |

| | | | | | | |
|------|------|------|-------|---|----------------------|-------------------|
| C 28 | 15.6 | 70.9 | 327 | 4 | US-09-252-991A-5577 | Sequence 5577, Ap |
| C 29 | 15.6 | 70.9 | 730 | 2 | US-08-743-637B-11 | Sequence 11, Appl |
| C 30 | 15.6 | 70.9 | 730 | 3 | US-08-526-840B-11 | Sequence 11, Appl |
| C 31 | 15.6 | 70.9 | 912 | 4 | US-09-252-991A-5679 | Sequence 5679, Ap |
| C 32 | 15.6 | 70.9 | 1007 | 3 | US-08-836-500A-1 | Sequence 1, Appl |
| C 33 | 15.6 | 70.9 | 1008 | 3 | US-08-721-979A-13 | Sequence 13, Appl |
| C 34 | 15.6 | 70.9 | 1008 | 4 | US-09-654-289-13 | Sequence 13, Appl |
| C 35 | 15.6 | 70.9 | 1008 | 4 | US-09-582-876-13 | Sequence 13, Appl |
| C 36 | 15.2 | 69.1 | 80 | 3 | US-09-039-555B-4 | Sequence 4, Appl |
| C 37 | 15.2 | 69.1 | 420 | 4 | US-09-252-991A-10335 | Sequence 10335, A |
| C 38 | 15.2 | 69.1 | 894 | 4 | US-09-252-991A-11338 | Sequence 11338, A |
| C 39 | 15.2 | 69.1 | 1335 | 4 | US-09-252-991A-11357 | Sequence 11357, A |
| C 40 | 15.2 | 69.1 | 1506 | 4 | US-09-252-991A-10637 | Sequence 10637, A |
| C 41 | 15.2 | 69.1 | 1605 | 4 | US-09-252-991A-10729 | Sequence 10729, A |
| C 42 | 15.2 | 69.1 | 4284 | 4 | US-09-252-991A-10434 | Sequence 10434, A |
| C 43 | 15.2 | 69.1 | 9704 | 4 | US-09-814-951A-3 | Sequence 3, Appl |
| C 44 | 15.2 | 69.1 | 44377 | 2 | US-08-804-227C-7 | Sequence 7, Appl |
| C 45 | 15.2 | 69.1 | 44377 | 2 | US-08-804-198-1 | Sequence 1, Appl |

ALIGNMENTS

RESULT 1
US-08-327-832-12
; Sequence 12, Application US/08327832
; Patent No. 5840832
; GENERAL INFORMATION:
; APPLICANT: Ono, Santa J.
; APPLICANT: Strominger, Jack L.
; TITLE OF INVENTION: Transcription Factor Regulating MHC
; TITLE OF INVENTION: Expression, cDNA and Genomic Clones Encoding Same and
; TITLE OF INVENTION: Retroviral Expression Constructs Thereof
; NUMBER OF SEQUENCES: 16
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Banner, Birch, McKie & Beckett
; STREET: 1001 G Street, N.W.
; CITY: Washington, D.C.
; STATE: District of Columbia
; COUNTRY: U.S.A.
; ZIP: 20001
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/327,832
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: Posorske, Laurence H.
; REGISTRATION NUMBER: 34,698
; REFERENCE/DOCKET NUMBER: 1107.46362
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-508-9153
; TELEFAX: 202-508-9299
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 22 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; ORGANISM: homo sapiens
US-08-327-832-12

Query Match 100.0%; Score 22; DB 2; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.46;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTCGATCGGGCGGCGGAGC 22

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Db      1  ATTCGATCGGGCGGGCGGCGAGC 22
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RESULT 2
US-08-828-584-12
; Sequence 12, Application US/08828584
; Patent No. 5908762
; GENERAL INFORMATION:
; APPLICANT: Ono, Santa J.
; APPLICANT: Strominger, Jack L.
; TITLE OF INVENTION: Transcription Factor Regulating MHC
; TITLE OF INVENTION: Expression, cDNA and Genomic Clones Encoding Same and
; TELECOMMUNICATION INFORMATION: Retroviral Expression Contracts Thereof
; NUMBER OF SEQUENCES: 16
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Banner, Birch, McKie & Beckett
; STREET: 1001 G Street, N.W.
; CITY: Washington, D.C.
; STATE: District of Columbia
; COUNTRY: U.S.A.
; ZIP: 20001
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/828,584
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Posorske, Laurence H.
; REGISTRATION NUMBER: 34,698
; REFERENCE/DOCKET NUMBER: 1107.46362
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 20-2 508-9153
; TELEFAX: 202 508-9299
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 22 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; ORGANISM: homo sapiens
; US-08-828-584-12
Query Match 100.0%; Score 22; DB 2; Length 22;
Best Local Similarity 100.0%; Pred. No. 0.46;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  ATTCGATCGGGCGGGCGGCGAGC 22
|||||
Db      1  ATTCGATCGGGCGGGCGGCGAGC 22
|||||
RESULT 3
PCT-US94-05659-16
; Sequence 16, Application PC/TUS9405659
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: TNF RESPONSIVE ELEMENT, TNF-INDUCED DNA-BINDING
; CORRESPONDENCE ADDRESS: 24
; ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02173
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/122,433
; FILING DATE: 22-SEP-1993
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/687,337
; FILING DATE: 18-APR-1991
; NAME: Reiter, Stephen E.
; REGISTRATION NUMBER: 31,192
; REFERENCE/DOCKET NUMBER: P31 9308
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-546-9392
; TELEFAX: 619-546-9392
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 46 base pairs
; QUERY MATCH 100.0%; Score 22; DB 5; Length 22;
; BEST LOCAL SIMILARITY 100.0%; PRED. NO. 0.46;
; MATCHES 22; CONSERVATIVE 0; MISMATCHES 0; INDELS 0; GAPS 0;

QY      1  ATTCGATCGGGCGGGCGGCGAGC 22
|||||
Db      1  ATTCGATCGGGCGGGCGGCGAGC 22
|||||
RESULT 4
US-08-122-433-12
; Sequence 12, Application US/08122433
; Patent No. 5683985
; GENERAL INFORMATION:
; APPLICANT: Chu, Barbara C.F.
; APPLICANT: Orgel, Leslie
; TITLE OF INVENTION: OLIGODEOXYNUCLEOTIDES AND
; TITLE OF INVENTION: OLIGONUCLEOTIDES USEFUL AS DECOYS FOR PROTEINS WHICH
; TITLE OF INVENTION: SELECTIVELY BIND TO DEFINED DNA SEQUENCES
; NUMBER OF SEQUENCES: 47
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PRETTY, SCHROEDER, BRUEGEMANN & CLARK
; STREET: 444 South Flower Street, Suite 2000
; CITY: Los Angeles
; STATE: California
; COUNTRY: USA
; ZIP: 90071
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/122,433
; FILING DATE: 22-SEP-1993
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/687,337
; FILING DATE: 18-APR-1991
; NAME: Reiter, Stephen E.
; REGISTRATION NUMBER: 31,192
; REFERENCE/DOCKET NUMBER: P31 9308
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-546-9392
; TELEFAX: 619-546-9392
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 46 base pairs
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; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/122,433
; FILING DATE: 22-SEP-1993
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/687,337
; FILING DATE: 18-APR-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Reiter, Stephen E.
; REGISTRATION NUMBER: 31,192
; REFERENCE/DOCKET NUMBER: P31 9308
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-546-1995
; TELEFAX: 619-546-9392
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 46 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: circular
; MOLECULE TYPE: other nucleic acid
; US-08-122-433-13

Query Match      81.8%; Score 18; DB 1; Length 46;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGGCGAGC 22
Db 24 GATCGGGCGGGCGGCGAGC 7

RESULT 8
US-08-458-434A-4/c
; Sequence 4, Application US/08458434A
; Patent No. 6083690
; GENERAL INFORMATION:
; APPLICANT: Mundy M.D., Gregory R.
; APPLICANT: Gosh-Choudhury Ph.D., Nandini
; APPLICANT: Feng Ph.D., Jian Q.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; TITLE OF INVENTION: OSTEOGENIC AGENTS
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: James C. Weseman, Esq.
; STREET: 401 B. Street, Suite 1700
; CITY: San Diego
; STATE: CA
; COUNTRY: USA
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/458,434A
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Weseman, James C.
; REGISTRATION NUMBER: 30,507
; REFERENCE/DOCKET NUMBER: P00060US0
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 699-3604
; TELEFAX: 619-236-1048
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2875 base pairs
; TYPE: nucleic acid
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; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-458-434A-4

Query Match      76.4%; Score 16.8; DB 3; Length 2875;
Best Local Similarity 90.0%; Pred. No. 60;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGCGAGC 22
Db 1805 TGGAGCGGGCGGGCGGCGAGC 1786

RESULT 9
US-08-458-434A-6/c
; Sequence 6, Application US/08458434A
; Patent No. 6083690
; GENERAL INFORMATION:
; APPLICANT: Harris Ph.D., Stephen E.
; APPLICANT: Mundy M.D., Gregory R.
; APPLICANT: Gosh-Choudhury Ph.D., Nandini
; APPLICANT: Feng Ph.D., Jian Q.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; TITLE OF INVENTION: OSTEOGENIC AGENTS
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: James C. Weseman, Esq.
; STREET: 401 B. Street, Suite 1700
; CITY: San Diego
; STATE: CA
; COUNTRY: USA
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/458,434A
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Weseman, James C.
; REGISTRATION NUMBER: 30,507
; REFERENCE/DOCKET NUMBER: P00060US0
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 699-3604
; TELEFAX: 619-236-1048
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 15144 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-458-434A-6

Query Match      76.4%; Score 16.8; DB 3; Length 15144;
Best Local Similarity 90.0%; Pred. No. 56;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGCGAGC 22
Db 1805 TGGAGCGGGCGGGCGGCGAGC 1786

RESULT 10
US-09-591-383-5/c
; Sequence 5, Application US/09591383
; Patent No. RE37984
; GENERAL INFORMATION:
; APPLICANT: Jackle, Herbert
```

;; Tautz, Diethard
;; TITLE OF INVENTION: PROCESS FOR ANALYZING LENGTH
;; NUMBER OF SEQUENCES: 6
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: BIRCH, STEWART, KOLASCH & BIRCH
;; STREET: 301 N. Washington Street, P.O. Box 747
;; CITY: Falls Church
;; STATE: Virginia
;; COUNTRY: United States of America
;; ZIP: 22046
;;
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.25
;; CURRENT APPLICATION NUMBER: US/09/591,383
;; FILING DATE: 09-Jun-2000
;; CLASSIFICATION: <Unknown>
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 07/681,494
;; FILING DATE: 10-JUN-1991
;; APPLICATION NUMBER: DE P3834636.2
;; FILING DATE: 11-OCT-1988
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Svensson, Leonard R.
;; REGISTRATION NUMBER: 30,330
;; REFERENCE/DOCKET NUMBER: 147-122PCT
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 703-241-1300
;; TELEFAX: 703-241-2848
;; TELEX: 248345
;; INFORMATION FOR SEQ ID NO: 5:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 379 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: DNA (genomic)
;; SEQUENCE DESCRIPTION: SEQ ID NO: 5:
US-09-591-383-5
Query Match 74.5%; Score 16.4; DB 1; Length 379;
Best Local Similarity 94.4%; Pred. No. 96;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 5 GATCGGGCGGGCGGAGC 22
Db 64 GATTGGGGCGGGCGGAGC 47

RESULT 11
US-08-145-617-5/c
; Sequence 5, Application US/08145617
; Patent No. 5766847
; GENERAL INFORMATION:
; APPLICANT: Jackle, Herbert
; APPLICANT: Tautz, Diethard
; TITLE OF INVENTION: PROCESS FOR ANALYZING LENGTH
; NUMBER OF SEQUENCES: 6
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BIRCH, STEWART, KOLASCH & BIRCH
; STREET: 301 N. Washington Street, P.O. Box 747
; CITY: Falls Church
; STATE: Virginia
; COUNTRY: United States of America
; ZIP: 22046
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS

;; SOFTWARE: PatentIn Release #1.0, Version #1.25
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/145,617
;; FILING DATE:
;; CLASSIFICATION: 435
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 07/681,494
;; FILING DATE: 10-JUN-1991
;; APPLICATION NUMBER: DE P3834636.2
;; FILING DATE: 11-OCT-1988
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Svensson, Leonard R.
;; REGISTRATION NUMBER: 30,330
;; REFERENCE/DOCKET NUMBER: 147-122PCT
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 703-241-1300
;; TELEFAX: 703-241-2848
;; TELEX: 248345
;; INFORMATION FOR SEQ ID NO: 5:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 379 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: DNA (genomic)
;; US-08-145-617-5
Query Match 74.5%; Score 16.4; DB 1; Length 379;
Best Local Similarity 94.4%; Pred. No. 96;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 5 GATCGGGCGGGCGGAGC 22
Db 64 GATTGGGGCGGGCGGAGC 47

RESULT 12
US-09-345-236B-145
; Sequence 145, Application US/09345236B
; Patent No. 6521454
; GENERAL INFORMATION:
; APPLICANT: Becnel, James J.
; APPLICANT: Tuku, Fukuda
; APPLICANT: Moser, Bettina
; APPLICANT: Cockburn, Andrew
; APPLICANT: White, Susan E.
; APPLICANT: Undeen, Albert H.
; TITLE OF INVENTION: No. 6521454el Baculoviruses, Insecticidal
; FILE OF INVENTION: Compositions, and Methods for Control of Invertebrates
; FILE REFERENCE: 21042.0004
; CURRENT APPLICATION NUMBER: US/09/345,236B
; CURRENT FILING DATE: 1999-06-30
; NUMBER OF SEQ ID NOS: 148
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 145
; LENGTH: 3997
; TYPE: DNA
; ORGANISM: mosquito baculovirus
; US-09-345-236B-145
Query Match 73.6%; Score 16.2; DB 4; Length 3997;
Best Local Similarity 85.7%; Pred. No. 1.1e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGAG 21
Db 2827 ATTCGTTGGGGTGGGTCGAG 2847

RESULT 13
US-09-345-236B-146/c
; Sequence 146, Application US/09345236B
; Patent No. 6521454

```
; GENERAL INFORMATION:
; APPLICANT: Becnel, James J.
; APPLICANT: Tokuo, Fukuda
; APPLICANT: Moser, Bettina
; APPLICANT: Cockburn, Andrew
; APPLICANT: White, Susan E.
; APPLICANT: Undeen, Albert H.
; TITLE OF INVENTION: No. 6521454el Baculoviruses, Insecticidal
; TITLE OF INVENTION: Compositions, and Methods for Control of Invertebrates
; FILE REFERENCE: 21042.0004
; CURRENT APPLICATION NUMBER: US/09/345,236B
; CURRENT FILING DATE: 1999-06-30
; NUMBER OF SEQ ID NOS: 148
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 146
; LENGTH: 3997
; TYPE: DNA
; ORGANISM: mosquito baculovirus
US-09-345-236B-146

Query Match 73.6%; Score 16.2; DB 4; Length 3997;
Best Local Similarity 85.7%; Pred. No. 1.1e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGCGAG 21
Db 1171 ATTCGTTGGGTGCGGGTCGAG 1151

RESULT 14
US-09-068-506-1
; Sequence 1, Application US/09068506A
; Patent No. 6569618
; GENERAL INFORMATION:
; APPLICANT: YASUE, Hirofumi
; APPLICANT: YOSHIMURA, Kumanoto
; TITLE OF INVENTION: DIAGNOSIS OF DISEASES ASSOCIATED WITH CORONARY
; FILE REFERENCE: 0032-245P
; CURRENT APPLICATION NUMBER: US/09/068,506A
; CURRENT FILING DATE: 1998-07-10
; NUMBER OF SEQ ID NOS: 72
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 9208
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: nnnnnnnnn = Intervening sequences of introns
US-09-068-506-1

Query Match 73.6%; Score 16.2; DB 4; Length 9208;
Best Local Similarity 85.7%; Pred. No. 1e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 ATTCGATCGGGCGGGCGGCGAG 21
Db 1489 ATGGGATAGGGCGGGCGGCGAG 1509

RESULT 15
US-08-153-563-4/c
; Sequence 4, Application US/08153563
; Patent No. 5693506
; GENERAL INFORMATION:
; APPLICANT: Rodriguez, Raymond L.
; TITLE OF INVENTION: PROCESS FOR PROTEIN PRODUCTION
; NUMBER OF INVENTION: IN PLANTS
; NUMBER OF SEQUENCES: 7
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend Kourie and Crew
; STREET: Stuart Street Tower, One Market Plaza
; CITY: San Francisco
```

```
; STATE: California
; COUNTRY: US
; ZIP: 94105-1493
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: US/08/153,563
; APPLICATION NUMBER: US/08/153,563
; FILING DATE: 16-NOV-1993
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Weber, Kenneth A.
; REGISTRATION NUMBER: 31,677
; REFERENCE/DOCKET NUMBER: 2307E-515
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 543-9600
; TELEFAX: (415) 543-5043
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 31 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: 1..31
; OTHER INFORMATION: /standard_name= "31 bp RAmv3E"
US-08-153-563-4

Query Match 71.8%; Score 15.8; DB 1; Length 31;
Best Local Similarity 89.5%; Pred. No. 1.9e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGCGAGC 22
Db 23 CGATCGAGGCGGCGGCGAGC 5

RESULT 16
US-09-038-227-9/c
; Sequence 9, Application US/09038227
; Patent No. 5917029
; GENERAL INFORMATION:
; APPLICANT: Yu, Su-May
; TITLE OF INVENTION: SUGAR-RESPONSIVE ENHANCERS
; TITLE OF INVENTION: IN ALPHA-AMYLASE GENES
; NUMBER OF SEQUENCES: 53
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 225 Franklin Street
; CITY: Boston
; STATE: MA
; COUNTRY: US
; ZIP: 02110-2804
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows95
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/038,227
; FILING DATE: 11-MAR-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Tsao, Y. Rocky
; REGISTRATION NUMBER: 34,053
; REFERENCE/DOCKET NUMBER: 05228/031001
; TELECOMMUNICATION INFORMATION:
```


APPLICANT: Corley, Neil C.
TITLE OF INVENTION: HUMAN REGULATORY MOLECULES
NUMBER OF SEQUENCES: 98
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/933,750C
FILING DATE: September 23, 1997
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PP-0356 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 97:
SEQUENCE CHARACTERISTICS:
LENGTH: 1112 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: TESTNOT07
CLONE: 3217567
US-08-933-750C-97

Query Match 71.8%; Score 15.8; DB 2; Length 1112;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAGC 22
Db 147 CGAGCGGGCGGGCGGCGGC 165

RESULT 20
US-09-234-613-97
Sequence 97, Application US/09234613
Patent No. 6132973
GENERAL INFORMATION:
APPLICANT: Lal, Preeti
APPLICANT: Hillman, Jennifer L.
APPLICANT: Bandman, Olga
APPLICANT: Shah, Purvi
APPLICANT: Au-Young, Janice
APPLICANT: Yue, Henry
APPLICANT: Guegler, Karl J.
APPLICANT: Corley, Neil C.
TITLE OF INVENTION: HUMAN REGULATORY MOLECULES
NUMBER OF SEQUENCES: 98
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/234,613
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/933,750
FILING DATE: September 23, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PP-0356 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
TELEX:
INFORMATION FOR SEQ ID NO: 97:
SEQUENCE CHARACTERISTICS:
LENGTH: 1112 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: TESTNOT07
CLONE: 3217567
US-09-234-613-97

Query Match 71.8%; Score 15.8; DB 3; Length 1112;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAGC 22
Db 147 CGAGCGGGCGGGCGGCGGC 165

RESULT 21
US-09-690-454-39
Sequence 39, Application US/09690454
Patent No. 6531447
GENERAL INFORMATION:
APPLICANT: Steven M. Ruben, et al.
TITLE OF INVENTION: 32 Human Secreted Proteins
FILE REFERENCE: P2006P1
CURRENT APPLICATION NUMBER: US/09/690,454
CURRENT FILING DATE: 2000-10-18
PRIOR APPLICATION NUMBER: 09/189,144
PRIOR FILING DATE: 1998-11-10
PRIOR APPLICATION NUMBER: 60/044,039
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/048,093
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/048,190
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/050,935
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/048,101
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/048,356
PRIOR FILING DATE: May 30, 1997
PRIOR APPLICATION NUMBER: 60/056,250
PRIOR FILING DATE: August 29, 1997
PRIOR APPLICATION NUMBER: 60/056,296
PRIOR FILING DATE: August 29, 1997
PRIOR APPLICATION NUMBER: 60/056,293
PRIOR FILING DATE: August 29, 1997
NUMBER OF SEQ ID NOS: 229
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 39
LENGTH: 1114

TYPE: DNA
ORGANISM: Homo sapiens
US-09-690-454-39

Query Match 71.8%; Score 15.8; DB 4; Length 1114;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 CGATCGGGGGCGGGCGGAG 22
DB 96 CGAGCGGGGGCGGGCGGCG 114

RESULT 22

US-07-991-587A-1/c
Sequence 1, Application US/07991587A
Patent No. 5384249
GENERAL INFORMATION:
APPLICANT: Sasaki, Katsutoshi
APPLICANT: Watanabe, Etsuyo
APPLICANT: Nishi, Tatsunari
APPLICANT: Sekine, Susumu
APPLICANT: Hanai, No. 5384249uo
APPLICANT: Hasegawa, Mamoru
TITLE OF INVENTION: '2 3 Sialyltransferase
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fitzpatrick, Cella, Harper & Scinto
STREET: 277 Park Avenue
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10172
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.44mb
COMPUTER: IBM PC
OPERATING SYSTEM: Dos 3.3
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/991,587A
FILING DATE: 19930526
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: JP-333661/1991
APPLICATION NUMBER: JP-091044/1992
FILING DATE: 17-12-1991
FILING DATE: 10-04-1992
ATTORNEY/AGENT INFORMATION:
NAME: Lawrence S. Perry
REGISTRATION NUMBER: 31,865
REFERENCE/DOCKET NUMBER: 1580.2
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-758-2400
TELEFAX: 212-758-2982
TELEX: 236262
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1919
TYPE: NUCLEIC ACID
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: cdna to mRNA
ORIGINAL SOURCE:
ORGANISM: human
CELL LINE: TYH cell
CELL TYPE: histiocyte cell
US-07-991-587A-1

Query Match 71.8%; Score 15.8; DB 1; Length 1919;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TCGATCGGGGGCGGGCGGAG 21

DB 59 TGGAGCGGGCGGGCGGAG 41

RESULT 23
US-08-309-985-1/c
Sequence 1, Application US/08309985
Patent No. 5494790
GENERAL INFORMATION:
APPLICANT: Sasaki, Katsutoshi
APPLICANT: Watanabe, Etsuyo
APPLICANT: Nishi, Tatsunari
APPLICANT: Sekine, Susumu
APPLICANT: Hanai, No. 5494790uo
APPLICANT: Hasegawa, Mamoru
TITLE OF INVENTION: '2 3 Sialyltransferase
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fitzpatrick, Cella, Harper & Scinto
STREET: 277 Park Avenue
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10172
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.44mb
COMPUTER: IBM PC
OPERATING SYSTEM: Dos 3.3
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/309,985
FILING DATE: 20-SEP-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/991,587
FILING DATE: 16-12-1992
APPLICATION NUMBER: JP-333661/1991
APPLICATION NUMBER: JP-091044/1992
FILING DATE: 17-12-1991
FILING DATE: 10-04-1992
ATTORNEY/AGENT INFORMATION:
NAME: Lawrence S. Perry
REGISTRATION NUMBER: 31,865
REFERENCE/DOCKET NUMBER: 1580.2
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-758-2400
TELEFAX: 212-758-2982
TELEX: 236262
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1919
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: cdna to mRNA
ORIGINAL SOURCE:
ORGANISM: human
CELL LINE: TYH cell
CELL TYPE: histiocyte cell
US-08-309-985-1

Query Match 71.8%; Score 15.8; DB 1; Length 1919;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 TCGATCGGGGGCGGGCGGAG 21
DB 59 TGGAGCGGGCGGGCGGAG 41

RESULT 24
US-07-973-324A-5/c
Sequence 5, Application US/07973324A

Patent No. 5460952
GENERAL INFORMATION:
APPLICANT: Yu, Su-May
TITLE OF INVENTION: Gene Expression System Comprising the
Promoter Region of the Alpha-Amylase Genes
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
STREET: 6300 Sears Tower, 233 South Wacker Drive
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/973.324A
FILING DATE: 04-NOV-1992
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Borun, Michael F.
REGISTRATION NUMBER: 25447
REFERENCE/DOCKET NUMBER: 31149
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 3314 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORGANISM: Rice (Oryzae sativa)
STRAIN: CV. M202
IMMEDIATE SOURCE:
LIBRARY: (EMBL) genomic
CLONE: '-Amy8-C
FEATURE:
NAME/KEY: CDS
LOCATION: join(1152..1241, 1385..2323, 2409..2690)
FEATURE:
NAME/KEY: mat peptide
LOCATION: join(1227..1241, 1385..2323, 2409..2690)
PUBLICATION INFORMATION:
AUTHORS: Yu et al., Su-May
TITLE: Regulation of '-amy1ase-encoding gene expression
TITLE: in germinating seeds and cultured cells of rice
JOURNAL: Gene
VOLUME: in press
US-07-973-324A-5

Query Match 71.8%; Score 15.8; DB 1; Length 3314;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGAGC 22
||||| ||||| ||||| ||||| |||||
Db 782 CGATCGAGGCGGCGGAGC 764

RESULT 25
US-08-343-380-5/c
Sequence 5, Application US/08343380
Patent No. 5712112
GENERAL INFORMATION:
APPLICANT: Yu, Su-May

APPLICANT: Liu, Li-Pei
TITLE OF INVENTION: Gene Expression System Comprising the
Promoter Region of the Alpha-Amylase Genes
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
STREET: 6300 Sears Tower, 233 South Wacker Drive
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/343.380
FILING DATE: 22-NOV-1994
CLASSIFICATION: 435
APPLICANT NUMBER: US 07/973.324
FILING DATE: 04-NOV-1992
ATTORNEY/AGENT INFORMATION:
NAME: Borun, Michael F.
REGISTRATION NUMBER: 25447
REFERENCE/DOCKET NUMBER: 31149
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 3314 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORGANISM: Rice (Oryzae sativa)
STRAIN: CV. M202
IMMEDIATE SOURCE:
LIBRARY: (EMBL) genomic
CLONE: '-Amy8-C
FEATURE:
NAME/KEY: CDS
LOCATION: join(1152..1241, 1385..2323, 2409..2690)
FEATURE:
NAME/KEY: mat peptide
LOCATION: join(1227..1241, 1385..2323, 2409..2690)
PUBLICATION INFORMATION:
AUTHORS: Yu et al., Su-May
TITLE: Regulation of '-amy1ase-encoding gene expression
TITLE: in germinating seeds and cultured cells of rice
JOURNAL: Gene
VOLUME: in press
US-08-343-380-5

Query Match 71.8%; Score 15.8; DB 1; Length 3314;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGAGC 22
||||| ||||| ||||| ||||| |||||
Db 782 CGATCGAGGCGGCGGAGC 764

RESULT 26
US-09-072-435-5/c
Sequence 5, Application US/09072435
Patent No. 6215051
GENERAL INFORMATION:
APPLICANT: Yu, Su-May

APPLICANT: Liu, Li-Fei
APPLICANT: Chan, Ming-Tsair
TITLE OF INVENTION: GENE EXPRESSION SYSTEM COMPRISING THE
TITLE OF INVENTION: PROMOTER REGION OF THE ALPHA-AMYLASE GENES
NUMBER OF SEQUENCES: 13
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
STREET: 233 South Wacker Drive/6300 Sears Tower
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/072,435
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/639,792
FILING DATE: 29-APR-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/973,324
FILING DATE: 04-NOV-1992
ATTORNEY/AGENT INFORMATION:
NAME: Gass, David A.
REGISTRATION NUMBER: 38,153
REFERENCE/DOCKET NUMBER: 28123/34274
TELECOMMUNICATION INFORMATION:
TELEPHONE: (312) 474-6300
TELEFAX: (312) 474-0448
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 3314 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Rice (Oryzae sativa)
STRAIN: CV. M202
IMMEDIATE SOURCE:
LIBRARY: (EMBL) genomic
CLONE: -Amy8-C
FEATURE:
NAME/KEY: CDS
LOCATION: join(1152..1241, 1385..2323, 2409..2690)
FEATURE:
NAME/KEY: mat_peptide
LOCATION: join(1227..1241, 1385..2323, 2409..2690)
PUBLICATION INFORMATION:
AUTHORS: Yu et al., Su-May
TITLE: Regulation of -amylase-encoding gene expression
TITLE: in germinating seeds and cultured cells of rice
JOURNAL: Gene
VOLUME: in press
US-09-072-435-5

Query Match 71.8%; Score 15.8; DB 3; Length 3314;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
Db 782 CGATCGAGGCGGGCGGAGC 764

RESULT 27

US-09-072-917A-5/c
; Sequence 5, Application US/09072917A

Patent No. 6288302
GENERAL INFORMATION:
APPLICANT: Yu, Su-May
APPLICANT: Liu, Li-Fei
APPLICANT: Chan, Ming-Tsair
TITLE OF INVENTION: Application of Alpha-Amylase Gene
TITLE OF INVENTION: Promoter and Signal Sequence in the Production of
Patent No. 6288302
TITLE OF INVENTION: Recombinant Proteins in Transgenic Plants and Transgenic
TITLE OF INVENTION: Plant Seeds
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
STREET: 233 South Wacker Drive/6300 Sears Tower
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/072,917A
FILING DATE:
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/509,962
FILING DATE: 01-AUG-1995
ATTORNEY/AGENT INFORMATION:
NAME: Gass, David A.
REGISTRATION NUMBER: 38,153
REFERENCE/DOCKET NUMBER: 28123/34257
TELECOMMUNICATION INFORMATION:
TELEPHONE: (312) 474-6300
TELEFAX: (312) 474-0448
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 3314 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: Rice (Oryzae sativa)
STRAIN: CV. M202
IMMEDIATE SOURCE:
LIBRARY: (EMBL) genomic
CLONE: alpha-Amy8-C
FEATURE:
NAME/KEY: CDS
LOCATION: join(1152..1241, 1385..2323, 2409..2690)
FEATURE:
NAME/KEY: mat_peptide
LOCATION: join(1227..1241, 1385..2323, 2409..2690)
PUBLICATION INFORMATION:
AUTHORS: Yu et al., Su-May
TITLE: Regulation of alpha-amylase-encoding gene expression
TITLE: in germinating seeds and cultured cells of rice
JOURNAL: Gene
VOLUME: in press
US-09-072-917A-5

Query Match 71.8%; Score 15.8; DB 3; Length 3314;
Best Local Similarity 89.5%; Pred. No. 1.6e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
Db 782 CGATCGAGGCGGGCGGAGC 764

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RESULT 28
US-09-252-991A-5577/c
; Sequence 5577, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; PRIOR FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 5577
; LENGTH: 327
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-5577

Query Match          70.9%; Score 15.6; DB 4; Length 327;
Best Local Similarity 81.8%; Pred. No. 2.1e-02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY      1  ATTCGATCGGGCGGCGGAGC 22
Db      282 ATTCGAACGTGGCGGCGGACC 261

RESULT 29
US-08-743-637B-11/c
; Sequence 11, Application US/08743637B
; Patent No. 5994066
; GENERAL INFORMATION:
; APPLICANT: BERGERON, Michel G.
; APPLICANT: PICARD, Francois J.
; APPLICANT: OUELLETTE, Marc
; APPLICANT: ROY, Paul H.
; TITLE OF INVENTION: SPECIES-SPECIFIC AND UNIVERSAL DNA
; TITLE OF INVENTION: PROBES AND AMPLIFICATION PRIMERS TO RAPIDLY DETECT AND
; TITLE OF INVENTION: IDENTIFY COMMON BACTERIAL PATHOGENS AND ASSOCIATED
; TITLE OF INVENTION: ANTIBIOTIC RESISTANCE GENES FROM CLINICAL SPECIMENS ...
; NUMBER OF SEQUENCES: 273
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: QUARLES & BRADY
; STREET: 411 EAST WISCONSIN AVENUE
; CITY: MILWAUKEE
; STATE: WISCONSIN
; COUNTRY: USA
; ZIP: 53202-4497
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/743,637B
; FILING DATE: 04-NOV-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/526,840
; FILING DATE: 11-SEP-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: BAKER, Jean C.
; REGISTRATION NUMBER: 35,433
; REFERENCE/DOCKET NUMBER: 850586.90012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (414) 277-5000
; TELEFAX: (414) 277-5591
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 730 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; MOLECULE TYPE: DNA (genomic)
; ORIGINAL SOURCE:
; ORGANISM: Klebsiella pneumoniae
US-08-526-840B-11/c

Query Match          70.9%; Score 15.6; DB 3; Length 730;
Best Local Similarity 81.8%; Pred. No. 2e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

```

```

; LENGTH: 730 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; ORIGINAL SOURCE:
; ORGANISM: Klebsiella pneumoniae
US-08-743-637B-11

Query Match          70.9%; Score 15.6; DB 2; Length 730;
Best Local Similarity 81.8%; Pred. No. 2e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY      1  ATTCGATCGGGCGGCGGAGC 22
Db      686 AATCGATCAGGCGGCGGAGC 665

RESULT 30
US-08-526-840B-11/c
; Sequence 11, Application US/08526840B
; Patent No. 6001564
; GENERAL INFORMATION:
; APPLICANT: BERGERON, Michel G.
; APPLICANT: OUELLETTE, Marc
; APPLICANT: ROY, Paul H.
; TITLE OF INVENTION: SPECIFIC AND UNIVERSAL PROBES AND
; TITLE OF INVENTION: AMPLIFICATION PRIMERS TO RAPIDLY DETECT AND IDENTIFY
; TITLE OF INVENTION: COMMON BACTERIAL PATHOGENS AND ANTIBIOTIC RESISTANCE GENES
; TITLE OF INVENTION: FROM CLINICAL SPECIMENS FOR ROUTINE DIAGNOSIS IN ...
; NUMBER OF SEQUENCES: 177
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: QUARLES & BRADY
; STREET: 411 East Wisconsin Avenue
; CITY: Milwaukee
; STATE: Wisconsin
; COUNTRY: USA
; ZIP: 53202-4497
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/526,840B
; FILING DATE: 11-SEP-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/304,732
; FILING DATE: 12-SEP-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: BAKER, Jean C.
; REGISTRATION NUMBER: 35,433
; REFERENCE/DOCKET NUMBER: 850586.90012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (414) 277-5000
; TELEFAX: (414) 277-5591
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 730 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; MOLECULE TYPE: DNA (genomic)
; ORIGINAL SOURCE:
; ORGANISM: Klebsiella pneumoniae
US-08-526-840B-11

Query Match          70.9%; Score 15.6; DB 3; Length 730;
Best Local Similarity 81.8%; Pred. No. 2e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

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QY 1 ATTCGATCGGGCGGCGGAGC 22

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Db      686 AATCGATCAGCGCGGCGAGC 665
;
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1007 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..1007
; US-08-836-500A-1
;
; Query Match 70.9%; Score 15.6; DB 3; Length 1007;
; Best Local Similarity 81.8%; Pred. No. 2e+02;
; Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
;
; QY      1 ATTCGATCGGGCGGGCGGAGC 22
;
; Db      934 AATCGATCAGCGCGGCGAGC 913
;
; RESULT 33
; US-08-721-979A-13/c
; Sequence 13, Application US/08721979A
; Patent No. 6113911
; GENERAL INFORMATION:
; APPLICANT: Binz, Hans
; APPLICANT: N'Guyen, Ngoc Thien
; APPLICANT: Baussant, Thierry
; APPLICANT: Trudel, Michel
; TITLE OF INVENTION: PEPTIDE FRAGMENT OF RESPIRATORY
; TITLE OF INVENTION: SYNCYTIAL VIRUS PROTEIN G, IMMUNOGENIC AGENT, PHARMACEUTICAL
; TITLE OF INVENTION: COMPOSITION CONTAINING IT AND PREPARATION PROCESS
; NUMBER OF SEQUENCES: 75
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Gordon W. Hueschen
; STREET: 715 The "H" Bldg., 310 East Michigan
; CITY: Kalamazoo
; STATE: MI
; COUNTRY: USA
; ZIP: 49007
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
;
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/721.979A
; FILING DATE: October 4, 1996
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: FR 94 04009
; FILING DATE: 06-APR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Hueschen, Gordon W.
; REGISTRATION NUMBER: 16,157
; REFERENCE/DOCKET NUMBER: PFS7PCTUS/dln
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 616-382-0030
; TELEFAX: 616-382-2030
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1008 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..1008
; OTHER INFORMATION: /note= "name : P40"
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Db      686 AATCGATCAGCGCGGCGAGC 665
;
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1007 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..1007
; US-08-836-500A-1
;
; Query Match 70.9%; Score 15.6; DB 4; Length 912;
; Best Local Similarity 81.8%; Pred. No. 2e+02;
; Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
;
; QY      1 ATTCGATCGGGCGGGCGGAGC 22
;
; Db      165 ATTCGACGTCGCGCGGCGAGC 186
;
; RESULT 32
; US-08-836-500A-1/c
; Sequence 1, Application US/08836500A
; Patent No. 6197929
; GENERAL INFORMATION:
; APPLICANT: Binz, Hans
; APPLICANT: Baussant, Thierry
; APPLICANT: Haeuw, Jean-Francois
; APPLICANT: Nguyen Ngoc, Thien
; TITLE OF INVENTION: Carrier Protein Having an Adjuvant
; TITLE OF INVENTION: Effect, Immunogenic Complex Containing It, Process for
; TITLE OF INVENTION: Their Preparation, Nucleotide Sequence and Vaccines
; Patent No. 6197929
; NUMBER OF SEQUENCES: 8
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Rocky, Milnamow & Katz, Ltd.
; STREET: 180 N. Stetson, 2 Prudential Plaza, Suite
; CITY: Chicago
; STATE: Illinois
; COUNTRY: U.S.A.
; ZIP: 60601
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836.500A
; FILING DATE:
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: Katz, Martin L.
; REGISTRATION NUMBER: 25,011
; REFERENCE/DOCKET NUMBER: PIE1514P0180US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 312-616-5400
; TELEFAX: 312-616-5460
```

US-08-721-979A-13

Query Match 70.9%; Score 15.6; DB 3; Length 1008;
Best Local Similarity 81.8%; Pred. No. 2e+02; 4; Indels 0; Gaps 0;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 1 ATTCGATCGGGCGGCGGCGAGC 22
Db 934 AATCGATCAGGCGGCGGCGAGC 913

RESULT 34

US-09-654-289-13/c
; Sequence 13, Application US/09654289
; Patent No. 6410030
; GENERAL INFORMATION:
; APPLICANT: Binz, Hans
; APPLICANT: N'Guyen, Ngoc Thien
; APPLICANT: Baussant, Thierry
; APPLICANT: Trudel, Michel
; TITLE OF INVENTION: PEPTIDE FRAGMENT OF RESPIRATORY
; TITLE OF INVENTION: SYNCYTIAL VIRUS PROTEIN G, IMMUNOGENIC AGENT, PHARMACEUTICAL
; TITLE OF INVENTION: COMPOSITION CONTAINING IT AND PREPARATION PROCESS
; NUMBER OF SEQUENCES: 75
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Gordon W. Hueschen
; STREET: 715 The "H" Bldg., 310 East Michigan
; STREET: Avenue
; CITY: Kalamazoo
; STATE: MI
; COUNTRY: USA
; ZIP: 49007
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/654,289
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/721,979
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Hueschen, Gordon W.
; REGISTRATION NUMBER: 16,157
; REFERENCE/DOCKET NUMBER: PF57PCTUS/dln
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 616-382-0030
; TELEFAX: 616-382-2030
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1008 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..1008
; OTHER INFORMATION: /note= "name : P40"
US-09-654-289-13

Query Match 70.9%; Score 15.6; DB 4; Length 1008;
Best Local Similarity 81.8%; Pred. No. 2e+02; 4; Indels 0; Gaps 0;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 1 ATTCGATCGGGCGGCGGCGAGC 22
Db 934 AATCGATCAGGCGGCGGCGAGC 913

RESULT 35

US-09-582-876-13/c
; Sequence 13, Application US/09582876
; Patent No. 6537556
; GENERAL INFORMATION:
; APPLICANT: Binz, Hans
; APPLICANT: N'Guyen, Ngoc Thien
; APPLICANT: Baussant, Thierry
; APPLICANT: Trudel, Michel
; TITLE OF INVENTION: PEPTIDE FRAGMENT OF RESPIRATORY
; TITLE OF INVENTION: SYNCYTIAL VIRUS PROTEIN G, IMMUNOGENIC AGENT, PHARMACEUTICAL
; TITLE OF INVENTION: COMPOSITION CONTAINING IT AND PREPARATION PROCESS
; NUMBER OF SEQUENCES: 75
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Gordon W. Hueschen
; STREET: 715 The "H" Bldg., 310 East Michigan
; STREET: Avenue
; CITY: Kalamazoo
; STATE: MI
; COUNTRY: USA
; ZIP: 49007
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/582,876
; FILING DATE: 30-Jun-2000
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/721,979
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Hueschen, Gordon W.
; REGISTRATION NUMBER: 16,157
; REFERENCE/DOCKET NUMBER: PF57PCTUS/dln
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 616-382-0030
; TELEFAX: 616-382-2030
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1008 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..1008
; OTHER INFORMATION: /note= "name : P40"
US-09-582-876-13

Query Match 70.9%; Score 15.6; DB 4; Length 1008;
Best Local Similarity 81.8%; Pred. No. 2e+02; 4; Indels 0; Gaps 0;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
Qy 1 ATTCGATCGGGCGGCGGCGAGC 22
Db 934 AATCGATCAGGCGGCGGCGAGC 913

RESULT 36

US-09-039-555B-4/c
; Sequence 4, Application US/09039555B
; Patent No. 6033856
; GENERAL INFORMATION:
; APPLICANT: Koerner, Kathrin
; APPLICANT: Mueller, Rolf
; APPLICANT: Sadlacek, Hans-Harald

TITLE OF INVENTION: PROMOTER OF THE CDC25B GENE, ITS
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 3000 K Street, N.W., Suite 500
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/039,555B
FILING DATE: 16-MAR-1998
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: DE 19710643.9
FILING DATE: 14-MAR-1997
ATTORNEY/AGENT INFORMATION:
NAME: Best, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 016779/0131
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 80 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "oligonucleotide"
US-09-039-555B-4

Query Match 69.1%; Score 15.2; DB 3; Length 80;
Best Local Similarity 85.0%; Pred. No. 3.3e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 34 TCGATCGGGCGGGCGGCGC 15

RESULT 37
US-09-252-991A-10335/G
; Sequence 10335, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 10335
; LENGTH: 420
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-10335

Query Match 69.1%; Score 15.2; DB 4; Length 420;
Best Local Similarity 85.0%; Pred. No. 3.1e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 243 TCGGTCGCGCGGGCGGATC 224

RESULT 38
US-09-252-991A-11338
; Sequence 11338, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11338
; LENGTH: 894
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11338

Query Match 69.1%; Score 15.2; DB 4; Length 894;
Best Local Similarity 85.0%; Pred. No. 3e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 675 TCGATCGGGCGGGCGGAGC 694

RESULT 39
US-09-252-991A-11357
; Sequence 11357, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11357
; LENGTH: 1335
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11357

Query Match 69.1%; Score 15.2; DB 4; Length 1335;
Best Local Similarity 85.0%; Pred. No. 2.9e+02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
DB 1046 TCGATCGGGCGGGCGGAGC 1065

RESULT 40
US-09-252-991A-10637
; Sequence 10637, Application US/09252991A
; Patent No. 6551795

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; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 10637
; LENGTH: 1506
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-10637

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```

Query Match      69.1%; Score 15.2; DB 4; Length 1506;
Best Local Similarity 85.0%; Pred. No. 2.9e-02;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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QY      3 TCGATCGGGCGGGCGGAGC 22
      ||| ||| ||| ||| ||| ||| |||
Db      435 TCGTTCGGCGGGCGGATC 454

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Search completed: February 18, 2004, 16:16:35
Job time : 32 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 16:16:06 ; Search time 195 Seconds
(without alignments)
415.589 Million cell updates/sec

Title: US-10-026-341A-2

Perfect score: 22

Sequence: 1 attcgatcgggcgggcgagc 22

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2449703 seqs, 1841816367 residues

Total number of hits satisfying chosen parameters: 4899406

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:

- 1: /cgn2_6/ptodata/1/pubna/US07_PUBCOMB.seq*
- 2: /cgn2_6/ptodata/1/pubna/PCT_NEW_PUB.seq*
- 3: /cgn2_6/ptodata/1/pubna/US06_NEW_PUB.seq*
- 4: /cgn2_6/ptodata/1/pubna/US06_PUBCOMB.seq*
- 5: /cgn2_6/ptodata/1/pubna/US07_NEW_PUB.seq*
- 6: /cgn2_6/ptodata/1/pubna/PCTUS_PUBCOMB.seq*
- 7: /cgn2_6/ptodata/1/pubna/US08_NEW_PUB.seq*
- 8: /cgn2_6/ptodata/1/pubna/US08_PUBCOMB.seq*
- 9: /cgn2_6/ptodata/1/pubna/US09A_PUBCOMB.seq*
- 10: /cgn2_6/ptodata/1/pubna/US09B_PUBCOMB.seq*
- 11: /cgn2_6/ptodata/1/pubna/US09C_PUBCOMB.seq*
- 12: /cgn2_6/ptodata/1/pubna/US09_NEW_PUB.seq*
- 13: /cgn2_6/ptodata/1/pubna/US09_NEW_PUB.seq*
- 14: /cgn2_6/ptodata/1/pubna/US10A_PUBCOMB.seq*
- 15: /cgn2_6/ptodata/1/pubna/US10B_PUBCOMB.seq*
- 16: /cgn2_6/ptodata/1/pubna/US10_NEW_PUB.seq*
- 17: /cgn2_6/ptodata/1/pubna/US60_NEW_PUB.seq*
- 18: /cgn2_6/ptodata/1/pubna/US60_PUBCOMB.seq*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----|-------------------|
| 1 | 22 | 100.0 | 22 | 9 | US-09-754-949-10 |
| 2 | 22 | 100.0 | 22 | 9 | US-09-918-889-20 |
| 3 | 22 | 100.0 | 22 | 9 | US-09-919-042-20 |
| 4 | 22 | 100.0 | 22 | 13 | US-10-417-422-11 |
| 5 | 22 | 100.0 | 22 | 14 | US-10-026-341A-2 |
| 6 | 21 | 95.5 | 21 | 11 | US-09-877-705A-81 |
| 7 | 21 | 95.5 | 21 | 11 | US-09-877-705A-82 |
| 8 | 21 | 95.5 | 21 | 11 | US-09-877-738A-81 |
| 9 | 21 | 95.5 | 21 | 11 | US-09-877-738A-82 |
| 10 | 21 | 95.5 | 21 | 11 | US-09-888-326-115 |
| 11 | 21 | 95.5 | 21 | 11 | US-09-888-326-229 |
| 12 | 21 | 95.5 | 21 | 11 | US-09-776-479-825 |
| 13 | 21 | 95.5 | 21 | 11 | US-09-776-479-826 |
| 14 | 21 | 95.5 | 21 | 15 | US-10-112-653-796 |
| 15 | 21 | 95.5 | 21 | 15 | US-10-112-653-797 |

Sequence 825, App
Sequence 826, App
Sequence 149, App
Sequence 149, App
Sequence 43, Appl
Sequence 43, Appl
Sequence 104, App
Sequence 104, App
Sequence 101, App
Sequence 101, App
Sequence 1661, App
Sequence 49, Appl
Sequence 1369, App
Sequence 248985,
Sequence 248985,
Sequence 248985,
Sequence 248986,
Sequence 47, Appl
Sequence 68, Appl
Sequence 1912, App
Sequence 2414, App
Sequence 3, Appl
Sequence 3, Appl
Sequence 11259, A
Sequence 1713, App
Sequence 1, Appl
Sequence 1, Appl
Sequence 1331, App
Sequence 231, App
Sequence 6, Appl

ALIGNMENTS

RESULT 1

US-09-754-949-10
; Sequence 10, Application US/09754949
; Patent No. US20020015939A1
; GENERAL INFORMATION:
; APPLICANT: MCCARTHY, JUSTIN
; APPLICANT: CORDELL, BARBARA
; TITLE OF INVENTION: METHODS FOR IDENTIFYING INHIBITORS OF
; TITLE OF INVENTION: NEURONAL DEGENERATION
; FILE REFERENCE: SCIOS.012A
; CURRENT APPLICATION NUMBER: US/09/754.949
; CURRENT FILING DATE: 2001-01-04
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 10
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-09-754-949-10

Query Match 100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGAGC 22
|||||
DB 1 ATTCGATCGGGCGGGCGAGC 22
|||||

RESULT 2

US-09-918-889-20
; Sequence 20, Application US/09918889
; Patent No. US20020053092A1
; GENERAL INFORMATION:
; APPLICANT: Readhead, Carol W.
; APPLICANT: Winston, Robert

APPLICANT: Koeffler, H. Phillip
APPLICANT: Muller, Carsten
TITLE OF INVENTION: NUCLEIC ACID CONSTRUCTS CONTAINING A
TITLE OF INVENTION: CYCLIN A1 PROMOTER, AND KIT
FILE REFERENCE: 18810-81603
CURRENT APPLICATION NUMBER: US/09/918,869
CURRENT FILING DATE: 2001-07-30
PRIOR APPLICATION NUMBER: US 09/292,723
PRIOR FILING DATE: 1999-04-15
PRIOR APPLICATION NUMBER: US 09/191,920
PRIOR FILING DATE: 1998-11-13
PRIOR APPLICATION NUMBER: US 60/065,825
PRIOR FILING DATE: 1997-11-14
PRIOR APPLICATION NUMBER: PCT/US98/24238
PRIOR FILING DATE: 1998-11-13
NUMBER OF SEQ ID NOS: 32
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 20
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Double-stranded oligonucleotide
US-09-918-869-20

Query Match 100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGCGAGC 22
DB 1 ATTCGATCGGGCGGGCGGCGAGC 22

RESULT 3
US-09-919-042-20
Sequence 20, Application US/09919042
Patent No. US20020056148A1
GENERAL INFORMATION:
APPLICANT: Readhead, Carol W.
APPLICANT: Winston, Robert
APPLICANT: Koeffler, H. Phillip
APPLICANT: Muller, Carsten
TITLE OF INVENTION: Transfection, Storage and Transfer of
TITLE OF INVENTION: Male Germ Cells for Generation of Selectable Transgenic Stem
FILE REFERENCE: 18810-81602
CURRENT APPLICATION NUMBER: US/09/919,042
CURRENT FILING DATE: 2001-07-30
PRIOR APPLICATION NUMBER: US 09/292,723
PRIOR FILING DATE: 1999-04-15
PRIOR APPLICATION NUMBER: US 09/191,920
PRIOR FILING DATE: 1998-11-13
PRIOR APPLICATION NUMBER: US 60/065,825
PRIOR FILING DATE: 1997-11-14
PRIOR APPLICATION NUMBER: PCT/US98/24238
PRIOR FILING DATE: 1998-11-13
NUMBER OF SEQ ID NOS: 32
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 20
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Double-stranded oligonucleotide
US-09-919-042-20

Query Match 100.0%; Score 22; DB 9; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGCGAGC 22
DB 1 ATTCGATCGGGCGGGCGGCGAGC 22

DB 1 ATTCGATCGGGCGGGCGGCGAGC 22

RESULT 4

US-10-417-422-11
Sequence 11, Application US/10417422
Publication No. US20030219720A1
GENERAL INFORMATION:
APPLICANT: MCCARTHY, JUSTIN
APPLICANT: CORDELL, BARBARA
APPLICANT: SCIOS, INC.
TITLE OF INVENTION: METHODS FOR IDENTIFYING INHIBITORS OF
TITLE OF INVENTION: NEURONAL DEGENERATION
FILE REFERENCE: SCIOS.012C1
CURRENT APPLICATION NUMBER: US/10/417,422
CURRENT FILING DATE: 2003-04-14
PRIOR APPLICATION NUMBER: 09/754949
PRIOR FILING DATE: 2001-02-04
PRIOR APPLICATION NUMBER: 60/175200
PRIOR FILING DATE: 2000-01-10
NUMBER OF SEQ ID NOS: 17
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 11
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Oligonucleotide
US-10-417-422-11

Query Match 100.0%; Score 22; DB 13; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGCGAGC 22
DB 1 ATTCGATCGGGCGGGCGGCGAGC 22

RESULT 5

US-10-026-341A-2
Sequence 2, Application US/10026341A
Publication No. US20020137715A1
GENERAL INFORMATION:
APPLICANT: Alain Mauviel
TITLE OF INVENTION: Blocking Sp1 Transcription Factor
TITLE OF INVENTION: Broadly Inhibits Extracellular Matrix Gene Expression In
TITLE OF INVENTION: Vitro and In Vivo: Implications for the Treatment of Tissue
TITLE OF INVENTION: Fibrosis
FILE REFERENCE: MAU01.NP001
CURRENT APPLICATION NUMBER: US/10/026,341A
CURRENT FILING DATE: 2001-12-21
PRIOR APPLICATION NUMBER: 60/259,585
PRIOR FILING DATE: 2001-01-03
NUMBER OF SEQ ID NOS: 3
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 2
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: synthetic oligonucleotide
US-10-026-341A-2

Query Match 100.0%; Score 22; DB 14; Length 22;
Best Local Similarity 100.0%; Pred. No. 1;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGCGAGC 22
DB 1 ATTCGATCGGGCGGGCGGCGAGC 22


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RESULT 6
US-09-877-705A-81
; Sequence 81, Application US/09877705A
; Publication No. US20030008283A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD FOR SCREENING FOR DRUG CANDIDATES FOR MODULATING TRANSCRIPTION FACTOR ACTIVITY
; FILE REFERENCE: 26757-704
; CURRENT APPLICATION NUMBER: US/09/877,705A
; CURRENT FILING DATE: 2001-08-16
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 81
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Transcription factor probe PP81
US-09-877-705A-81

Query Match          95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
    |||||
Db 1 ATTCGATCGGGCGGGCGGAG 21
    |||||

RESULT 7
US-09-877-705A-82/c
; Sequence 82, Application US/09877705A
; Publication No. US20030008283A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD FOR SCREENING FOR DRUG CANDIDATES FOR MODULATING TRANSCRIPTION FACTOR ACTIVITY
; FILE REFERENCE: 26757-704
; CURRENT APPLICATION NUMBER: US/09/877,705A
; CURRENT FILING DATE: 2001-08-16
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 82
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Transcription factor probe PP82
US-09-877-705A-82

Query Match          95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
    |||||
Db 1 ATTCGATCGGGCGGGCGGAG 21
    |||||

RESULT 8
US-09-877-738A-81
; Sequence 81, Application US/09877738A
; Publication No. US20030022173A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD AND KIT FOR ISOLATING DNA PROBES THAT BIND TO ACTIVATED
; FILE REFERENCE: 26757-701
; CURRENT APPLICATION NUMBER: US/09/877,738A
; CURRENT FILING DATE: 2001-06-01
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 81
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Transcription factor probe PP81
US-09-877-738A-81

Query Match          95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
    |||||
Db 1 ATTCGATCGGGCGGGCGGAG 21
    |||||

RESULT 9
US-09-877-738A-82/c
; Sequence 82, Application US/09877738A
; Publication No. US20030022173A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD AND KIT FOR ISOLATING DNA PROBES THAT BIND TO ACTIVATED
; FILE REFERENCE: 26757-701
; CURRENT APPLICATION NUMBER: US/09/877,738A
; CURRENT FILING DATE: 2001-06-01
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 82
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Transcription factor probe PP82
US-09-877-738A-82

Query Match          95.5%; Score 21; DB 11; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
    |||||
Db 21 ATTCGATCGGGCGGGCGGAG 1
    |||||

RESULT 10
US-09-888-326-115
; Sequence 115, Application US/09888326
; Publication No. US20030026801A1
; GENERAL INFORMATION:
; APPLICANT: Weiner, George
; APPLICANT: Hartmann, Gunther
; TITLE OF INVENTION: Methods for Enhancing Antibody-Induced
; FILE REFERENCE: C1039/7052 (AWS)
; CURRENT APPLICATION NUMBER: US/09/888,326
; CURRENT FILING DATE: 2001-06-22
; PRIOR APPLICATION NUMBER: US 60/213,346
; PRIOR FILING DATE: 2000-06-22
; NUMBER OF SEQ ID NOS: 848
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 115
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
; NAME/KEY: misc.feature
; LOCATION: (0)...(0)
; OTHER INFORMATION: phosphodiester backbone
US-09-888-326-115
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Query Match 95.5%; Score 21; DB 11; Length 21;
 Best Local Similarity 100.0%; Pred. No. 2.9;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
 |||||
 DB 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 11
 US-09-888-326-229/c
 ; Sequence 229, Application US/09888326
 ; Publication No. US20030026801A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Weiner, George
 ; APPLICANT: Hartmann, Gunther
 ; TITLE OF INVENTION: Methods for Enhancing Antibody-Induced
 ; TITLE OF INVENTION: Cell Lysis and Treating Cancer
 ; FILE REFERENCE: C1039/7052 (AWS)
 ; CURRENT APPLICATION NUMBER: US/09/888,326
 ; CURRENT FILING DATE: 2001-06-22
 ; PRIOR APPLICATION NUMBER: US 60/213,346
 ; PRIOR FILING DATE: 2000-06-22
 ; NUMBER OF SEQ ID NOS: 848
 ; SOFTWARE: FastSeq for Windows Version 3.0
 ; SEQ ID NO 229
 ; LENGTH: 21
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; NAME/KEY: misc_feature
 ; LOCATION: (0)...(0)
 ; OTHER INFORMATION: phosphodiester backbone
 US-09-888-326-229

Query Match 95.5%; Score 21; DB 11; Length 21;
 Best Local Similarity 100.0%; Pred. No. 2.9;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
 |||||
 DB 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 12
 US-09-776-479-825
 ; Sequence 825, Application US/09776479
 ; Publication No. US20030087848A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Bratzler, Robert L.
 ; APPLICANT: Petersen, Deanna M.
 ; APPLICANT: Fouron, Yves
 ; TITLE OF INVENTION: Immunostimulatory Nucleic Acids for the
 ; TITLE OF INVENTION: Treatment of Asthma and Allergy
 ; FILE REFERENCE: C1037/7013 (HCL/MAT)
 ; CURRENT APPLICATION NUMBER: US/09/776,479
 ; CURRENT FILING DATE: 2001-02-02
 ; PRIOR APPLICATION NUMBER: US 60/179,991
 ; PRIOR FILING DATE: 2000-02-03
 ; NUMBER OF SEQ ID NOS: 1093
 ; SOFTWARE: FastSeq for Windows Version 3.0
 ; SEQ ID NO 825
 ; LENGTH: 21
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Synthetic Sequence
 US-09-776-479-825

Query Match 95.5%; Score 21; DB 11; Length 21;
 Best Local Similarity 100.0%; Pred. No. 2.9;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATTCGATCGGGCGGGCGGAG 21
 |||||
 DB 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 13
 US-09-776-479-826/c
 ; Sequence 826, Application US/09776479
 ; Publication No. US20030087848A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Bratzler, Robert L.
 ; APPLICANT: Petersen, Deanna M.
 ; APPLICANT: Fouron, Yves
 ; TITLE OF INVENTION: Immunostimulatory Nucleic Acids for the
 ; TITLE OF INVENTION: Treatment of Asthma and Allergy
 ; FILE REFERENCE: C1037/7013 (HCL/MAT)
 ; CURRENT APPLICATION NUMBER: US/09/776,479
 ; CURRENT FILING DATE: 2001-02-02
 ; PRIOR APPLICATION NUMBER: US 60/179,991
 ; PRIOR FILING DATE: 2000-02-03
 ; NUMBER OF SEQ ID NOS: 1093
 ; SOFTWARE: FastSeq for Windows Version 3.0
 ; SEQ ID NO 826
 ; LENGTH: 21
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Synthetic Sequence
 US-09-776-479-826

Query Match 95.5%; Score 21; DB 11; Length 21;
 Best Local Similarity 100.0%; Pred. No. 2.9;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
 |||||
 DB 21 ATTCGATCGGGCGGGCGGAG 1

RESULT 14
 US-10-112-653-796
 ; Sequence 796, Application US/10112653
 ; Publication No. US20030050268A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Krieg, Arthur M.
 ; APPLICANT: Berg, Daniel J.
 ; TITLE OF INVENTION: IMMUNOSTIMULATORY NUCLEIC ACID FOR
 ; TITLE OF INVENTION: TREATMENT OF NON-ALLERGIC INFLAMMATORY DISEASES
 ; FILE REFERENCE: C01039/70060(AWS)
 ; CURRENT APPLICATION NUMBER: US/10/112,653
 ; CURRENT FILING DATE: 2002-03-29
 ; PRIOR APPLICATION NUMBER: US 60/279,642
 ; PRIOR FILING DATE: 2001-03-29
 ; NUMBER OF SEQ ID NOS: 1040
 ; SOFTWARE: FastSeq for Windows Version 3.0
 ; SEQ ID NO 796
 ; LENGTH: 21
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Synthetic Oligonucleotide
 US-10-112-653-796

Query Match 95.5%; Score 21; DB 15; Length 21;
 Best Local Similarity 100.0%; Pred. No. 2.9;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
 |||||
 DB 1 ATTCGATCGGGCGGGCGGAG 21

RESULT 15
US-10-112-653-797/c
; Sequence 797, Application US/10112653
; Publication No. US20030050268A1
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; TITLE OF INVENTION: IMMUNOSTIMULATORY NUCLEIC ACID FOR
; TITLE OF INVENTION: TREATMENT OF NON-ALLERGIC INFLAMMATORY DISEASES
; FILE REFERENCE: C01039/70060(AWS)
; CURRENT APPLICATION NUMBER: US/10/112,653
; CURRENT FILING DATE: 2002-03-29
; PRIOR APPLICATION NUMBER: US 60/279,642
; PRIOR FILING DATE: 2001-03-29
; NUMBER OF SEQ ID NOS: 1040
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 797
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-10-112-653-797

Query Match 95.5%; Score 21; DB 15; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTGATCGGGCGGGCGGAG 21
Db 21 ATTGATCGGGCGGGCGGAG 1

RESULT 16
US-10-017-995-825
; Sequence 825, Application US/10017995
; Publication No. US20030055014A1
; GENERAL INFORMATION:
; APPLICANT: Bratzler, Robert L.
; TITLE OF INVENTION: Inhibition of Angiogenesis by Nucleic Acids
; FILE REFERENCE: C1037/7025 (HCL/MAT)
; CURRENT APPLICATION NUMBER: US/10/017,995
; CURRENT FILING DATE: 2001-12-18
; PRIOR APPLICATION NUMBER: US 60/255,534
; PRIOR FILING DATE: 2000-12-14
; NUMBER OF SEQ ID NOS: 1093
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 825
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-10-017-995-825

Query Match 95.5%; Score 21; DB 15; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTGATCGGGCGGGCGGAG 21
Db 1 ATTGATCGGGCGGGCGGAG 21

RESULT 17
US-10-017-995-826/c
; Sequence 826, Application US/10017995
; Publication No. US20030055014A1
; GENERAL INFORMATION:
; APPLICANT: Bratzler, Robert L.
; TITLE OF INVENTION: Inhibition of Angiogenesis by Nucleic Acids
; FILE REFERENCE: C1037/7025 (HCL/MAT)

; CURRENT APPLICATION NUMBER: US/10/017,995
; CURRENT FILING DATE: 2001-12-18
; PRIOR APPLICATION NUMBER: US 60/255,534
; PRIOR FILING DATE: 2000-12-14
; NUMBER OF SEQ ID NOS: 1093
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 826
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-10-017-995-826

Query Match 95.5%; Score 21; DB 15; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTGATCGGGCGGGCGGAG 21
Db 21 ATTGATCGGGCGGGCGGAG 1

RESULT 18
US-09-877-705A-149/c
; Sequence 149, Application US/09877705A
; Publication No. US20030008283A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD FOR SCREENING FOR DRUG CANDIDATES FOR MODULATING TRANSCR
; TITLE OF INVENTION: FACTOR ACTIVITY
; FILE REFERENCE: 26757-704
; CURRENT APPLICATION NUMBER: US/09/877,705A
; CURRENT FILING DATE: 2001-08-16
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 149
; LENGTH: 63
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hybridization probe MP82
US-09-877-705A-149

Query Match 95.5%; Score 21; DB 11; Length 63;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTGATCGGGCGGGCGGAG 21
Db 63 ATTGATCGGGCGGGCGGAG 43

RESULT 19
US-09-877-738A-149/c
; Sequence 149, Application US/09877738A
; Publication No. US20030022173A1
; GENERAL INFORMATION:
; APPLICANT: Li, Jason
; TITLE OF INVENTION: METHOD AND KIT FOR ISOLATING DNA PROBES THAT BIND TO ACTIVATED
; TITLE OF INVENTION: TRANSCRIPTION FACTORS
; FILE REFERENCE: 26757-701
; CURRENT APPLICATION NUMBER: US/09/877,738A
; CURRENT FILING DATE: 2001-06-01
; NUMBER OF SEQ ID NOS: 162
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 149
; LENGTH: 63
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hybridization probe MP82
US-09-877-738A-149

Query Match 95.5%; Score 21; DB 11; Length 63;
Best Local Similarity 100.0%; Pred. No. 2.5; Indels 0; Gaps 0;
Matches 21; Conservative 0; Mismatches 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
|||||
DB 63 ATTCGATCGGGCGGGCGGAG 43

RESULT 20

US-10-437-107-43
; Sequence 43, Application US/10437107
; Publication No. US20030186313A1
; GENERAL INFORMATION:
; APPLICANT: Fuqua, Suzanne
; APPLICANT: Allred, D.
; APPLICANT: Hopp, Torsten A.
; APPLICANT: O'Connell, Peter
; TITLE OF INVENTION: Methods and Composition in Breast Cancer Diagnosis and Therapeutic
; FILE REFERENCE: P02102US2
; CURRENT FILING DATE: 2003-05-13
; PRIOR APPLICATION NUMBER: US/10/437,107
; PRIOR FILING DATE: 2002-01-18
; PRIOR APPLICATION NUMBER: US 60/262,990
; PRIOR FILING DATE: 2001-01-19
; PRIOR APPLICATION NUMBER: US 60/304,018
; PRIOR FILING DATE: 2001-07-09
; NUMBER OF SEQ ID NOS: 49
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 43
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Estrogen Response Element
US-10-437-107-43

Query Match 92.7%; Score 20.4; DB 13; Length 22;
Best Local Similarity 95.5%; Pred. No. 5.5;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 22
|||||
DB 1 ATTCGATCAGGGCGGGCGGAG 22

RESULT 21

US-10-052-092-43
; Sequence 43, Application US/10052092
; Publication No. US20030027778A1
; GENERAL INFORMATION:
; APPLICANT: Fuqua, Suzanne
; APPLICANT: Allred, D.
; APPLICANT: Hopp, Torsten A.
; APPLICANT: O'Connell, Peter
; TITLE OF INVENTION: Methods and Composition in Breast Cancer Diagnosis and Therapeutic
; FILE REFERENCE: P02102US2
; CURRENT APPLICATION NUMBER: US/10/052,092
; CURRENT FILING DATE: 2002-01-18
; PRIOR APPLICATION NUMBER: US 60/262,990
; PRIOR FILING DATE: 2001-01-19
; PRIOR APPLICATION NUMBER: US 60/304,018
; PRIOR FILING DATE: 2001-07-09
; NUMBER OF SEQ ID NOS: 49
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 43
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Estrogen Response Element
US-10-052-092-43

US-10-052-092-43

Query Match 92.7%; Score 20.4; DB 15; Length 22;
Best Local Similarity 95.5%; Pred. No. 5.5;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 22
|||||
DB 1 ATTCGATCAGGGCGGGCGGAG 22

RESULT 22

US-09-954-987B-104
; Sequence 104, Application US/09954987B
; Publication No. US20030104523A1
; GENERAL INFORMATION:
; APPLICANT: Stefan Bauer
; APPLICANT: Grayson B. Lipford
; APPLICANT: Hermann Wagner
; TITLE OF INVENTION: PROCESS FOR HIGH THROUGHPUT SCREENING OF
; TITLE OF INVENTION: CPG-BASED IMMUNO-AGONIST/ANTAGONIST
; FILE REFERENCE: C1041/2016 (AMS)
; CURRENT APPLICATION NUMBER: US/09/954,987B
; CURRENT FILING DATE: 2001-09-17
; PRIOR APPLICATION NUMBER: US 60/233,035
; PRIOR FILING DATE: 2000-09-15
; PRIOR APPLICATION NUMBER: US 60/263,657
; PRIOR FILING DATE: 2001-01-23
; PRIOR APPLICATION NUMBER: US 60/291,726
; PRIOR FILING DATE: 2001-05-17
; PRIOR APPLICATION NUMBER: US 60/300,210
; PRIOR FILING DATE: 2001-06-22
; NUMBER OF SEQ ID NOS: 230
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 104
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
US-09-954-987B-104

Query Match 90.9%; Score 20; DB 11; Length 20;
Best Local Similarity 100.0%; Pred. No. 8.4;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAG 22
|||||
DB 1 TCGATCGGGCGGGCGGAG 20

RESULT 23

US-10-265-072-102
; Sequence 102, Application US/10265072
; Publication No. US20030166001A1
; GENERAL INFORMATION:
; APPLICANT: Lipford, Grayson
; TITLE OF INVENTION: TOLL-LIKE RECEPTOR 3 SIGNALING AGONISTS AND ANTAGONISTS
; FILE REFERENCE: C01041.70031.US
; CURRENT APPLICATION NUMBER: US/10/265,072
; CURRENT FILING DATE: 2002-10-05
; NUMBER OF SEQ ID NOS: 117
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 102
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Synthetic oligonucleotide
US-10-265-072-102

Query Match 90.9%; Score 20; DB 13; Length 20;
Best Local Similarity 100.0%; Pred. No. 8.4;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 22
Db 1 TCGATCGGGCGGGCGGAGC 20

RESULT 24
US-10-145-289-10/c
; Sequence 10, Application US/10145289
; Publication No. US20030077812A1
; GENERAL INFORMATION:
; APPLICANT: James G. McArthur
; APPLICANT: Dale John Talbot
; APPLICANT: Andrew D. Simmons
; APPLICANT: Ryan McGuinness
; APPLICANT: Michael Kelly
; APPLICANT: Lisa V. Tsui
; APPLICANT: Thomas Dull
; TITLE OF INVENTION: LENTIVIRAL VECTORS ENCODING CLOTTING
; FILE REFERENCE: 131.2USU1
; CURRENT APPLICATION NUMBER: US/10/145,289
; PRIOR FILING DATE: 2002-05-14
; PRIOR APPLICATION NUMBER: 60/291,083
; PRIOR FILING DATE: 2001-05-14
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 10
; LENGTH: 183
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: enhancer element
US-10-145-289-10

Query Match 79.1%; Score 17.4; DB 15; Length 183;
Best Local Similarity 94.7%; Pred. No. 51;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
Db 183 CGATCGGGCGGGCGGAGC 165

RESULT 25
US-10-311-455-1661
; Sequence 1661, Application US/10311455
; Publication No. US20030143606A1
; GENERAL INFORMATION:
; APPLICANT: OLEK Alexander
; APPLICANT: PIEPENBROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with the Immune System by Determining Cytosine Methylation
; FILE REFERENCE: 5013.1014
; CURRENT APPLICATION NUMBER: US/10/311,455
; CURRENT FILING DATE: 2002-12-16
; PRIOR APPLICATION NUMBER: PCT/EP01/07537
; PRIOR FILING DATE: 2001-07-02
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 2424
; SEQ ID NO 1661
; LENGTH: 9646
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-311-455-1661

Query Match 79.1%; Score 17.4; DB 13; Length 9646;
Best Local Similarity 94.7%; Pred. No. 49;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAGC 21
Db 5203 TCGATCGGGCGGGCGGAGC 5221

RESULT 26
US-09-769-734-49/c
; Sequence 49, Application US/09769734
; Publication No. US20030143666A1
; GENERAL INFORMATION:
; APPLICANT: Ecopia BioSciences Inc.
; TITLE OF INVENTION: Genetic Locus for Everninomicin Biosynthesis
; FILE REFERENCE: PA 005-US
; CURRENT APPLICATION NUMBER: US/09/769,734
; CURRENT FILING DATE: 2001-01-26
; NUMBER OF SEQ ID NOS: 59
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 49
; LENGTH: 11115
; TYPE: DNA
; ORGANISM: M. carbonacea
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (8)..(1207)
; OTHER INFORMATION: ORF 41 (positive strandedness)
; OTHER INFORMATION: incomplete: C-terminus only
; NAME/KEY: misc feature
; LOCATION: (1213)..(2331)
; OTHER INFORMATION: ORF 42 (positive strandedness)
; NAME/KEY: misc feature
; LOCATION: (2364)..(3611)
; OTHER INFORMATION: ORF 43 (positive strandedness)
; NAME/KEY: misc feature
; LOCATION: (3623)..(4243)
; OTHER INFORMATION: ORF 44 (positive strandedness)
; NAME/KEY: misc feature
; LOCATION: (4149)..(5177)
; OTHER INFORMATION: ORF 45 (positive strandedness)
; NAME/KEY: misc feature
; LOCATION: (5177)..(6094)
; OTHER INFORMATION: ORF 46 (negative strandedness)
; NAME/KEY: misc feature
; LOCATION: (6271)..(7824)
; OTHER INFORMATION: ORF 47 (negative strandedness)
; NAME/KEY: misc feature
; LOCATION: (7903)..(8760)
; OTHER INFORMATION: ORF 48 (negative strandedness)
; NAME/KEY: misc feature
; LOCATION: (8781)..(9800)
; OTHER INFORMATION: ORF 49 (negative strandedness)
US-09-769-734-49

Query Match 78.2%; Score 17.2; DB 13; Length 11115;
Best Local Similarity 86.4%; Pred. No. 59;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 ATTGATCGGGCGGGCGGAGC 22
Db 6118 ATTGATCGGGCGGGCGGAGC 6097

RESULT 27
US-10-292-798-1369/c
; Sequence 1369, Application US/10292798
; Publication No. US20030235833A1
; GENERAL INFORMATION:
; APPLICANT: SUWA, MAKIKO
; APPLICANT: ASAI, KIYOSHI
; APPLICANT: AKIYAMA, YUTAKA

APPLICANT: ABRATANI, HIROYUKI
TITLE OF INVENTION: GUANOSINE TRIPHOSPHATE-BINDING PROTEIN COUPLED RECEPTORS
FILE REFERENCE: 084335/166
CURRENT APPLICATION NUMBER: US/10/292,798
CURRENT FILING DATE: 2002-11-13
PRIOR APPLICATION NUMBER: 10/017,161
PRIOR FILING DATE: 2001-12-18
PRIOR APPLICATION NUMBER: JP 2001-246789
PRIOR FILING DATE: 2001-06-18
NUMBER OF SEQ ID NOS: 2070
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 1369
LENGTH: 744802
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
LOCATION: source
FEATURE:
LOCATION: (1)..(744802)
FEATURE:
NAME/KEY: CDS
LOCATION: (201)..(246)
FEATURE:
NAME/KEY: CDS
LOCATION: (25640)..(25677)
FEATURE:
NAME/KEY: CDS
LOCATION: (27078)..(27094)
FEATURE:
NAME/KEY: CDS
LOCATION: (141192)..(141769)
FEATURE:
NAME/KEY: CDS
LOCATION: (159571)..(159606)
FEATURE:
NAME/KEY: CDS
LOCATION: (174525)..(174575)
FEATURE:
NAME/KEY: CDS
LOCATION: (234891)..(235013)
FEATURE:
NAME/KEY: CDS
LOCATION: (235514)..(235560)
FEATURE:
NAME/KEY: CDS
LOCATION: (279677)..(279729)
FEATURE:
NAME/KEY: CDS
LOCATION: (408660)..(409123)
FEATURE:
NAME/KEY: CDS
LOCATION: (409204)..(409669)
FEATURE:
NAME/KEY: CDS
LOCATION: (428381)..(428396)
FEATURE:
NAME/KEY: CDS
LOCATION: (472204)..(472330)
FEATURE:
NAME/KEY: CDS
LOCATION: (714252)..(714355)
FEATURE:
NAME/KEY: CDS
LOCATION: (714447)..(714529)
FEATURE:
NAME/KEY: CDS
LOCATION: (739794)..(739891)
FEATURE:
NAME/KEY: CDS
LOCATION: (744484)..(744602)
FEATURE:
NAME/KEY: modified_base
LOCATION: (51812)..(51911)

OTHER INFORMATION: a, t, c, g, unknown or other
FEATURE:
NAME/KEY: modified_base
LOCATION: (57122)..(57221)
OTHER INFORMATION: a, t, c, g, unknown or other
FEATURE:
NAME/KEY: modified_base
LOCATION: (79368)..(79467)
OTHER INFORMATION: a, t, c, g, unknown or other
FEATURE:
NAME/KEY: modified_base
LOCATION: (293951)..(294050)
OTHER INFORMATION: a, t, c, g, unknown or other
FEATURE:
NAME/KEY: modified_base
LOCATION: (310089)..(310188)
OTHER INFORMATION: a, t, c, g, unknown or other
FEATURE:
NAME/KEY: modified_base
LOCATION: (332935)..(332935)
OTHER INFORMATION: a, t, c, g, unknown or other
FEATURE:
NAME/KEY: modified_base
LOCATION: (332992)..(332992)
OTHER INFORMATION: a, t, c, g, unknown or other
FEATURE:
NAME/KEY: modified_base
LOCATION: (362002)..(362101)
OTHER INFORMATION: a, t, c, g, unknown or other
FEATURE:
NAME/KEY: modified_base
LOCATION: (639781)..(639880)
OTHER INFORMATION: a, t, c, g, unknown or other
US-10-292-798-1369

Query Match 78.2%; Score 17.2; DB 12; Length 744802;
Best Local Similarity 86.4%; Pred. No. 31;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAC 22
DB 384946 ATTCGATAGGGTGGGGCGTGC 384925

RESULT 28
US-10-027-632-248985
Sequence 248985, Application US/10027632
Publication No. US20030204075A9
GENERAL INFORMATION:
APPLICANT: Wang, David G.
TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
FILE REFERENCE: 108827.129
CURRENT APPLICATION NUMBER: US/10/027,632
CURRENT FILING DATE: 2002-04-30
PRIOR APPLICATION NUMBER: US 60/218,006
PRIOR FILING DATE: 2000-07-12
PRIOR APPLICATION NUMBER: US 60/198,676
PRIOR FILING DATE: 2000-04-20
PRIOR APPLICATION NUMBER: US 60/193,483
PRIOR FILING DATE: 2000-03-29
PRIOR APPLICATION NUMBER: US 60/185,218
PRIOR FILING DATE: 2000-02-24
PRIOR APPLICATION NUMBER: US 60/167,363
PRIOR FILING DATE: 1999-11-23
PRIOR APPLICATION NUMBER: US 60/156,358
PRIOR FILING DATE: 1999-09-28
PRIOR APPLICATION NUMBER: US 60/146,002
PRIOR FILING DATE: 1999-08-09
NUMBER OF SEQ ID NOS: 325720
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 248985
LENGTH: 1073

```
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-248985

Query Match      76.4%; Score 16.8; DB 13; Length 1073;
Best Local Similarity 90.0%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTCGATCGGGCGGGCGGAG 21
DB 922 TTCAATCGGGCGGGGGGAG 941

RESULT 29
US-10-027-632-248986
; Sequence 248986, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; TITLE OF INVENTION: Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 248986
; LENGTH: 1073
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-248986

Query Match      76.4%; Score 16.8; DB 13; Length 1073;
Best Local Similarity 90.0%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTCGATCGGGCGGGCGGAG 21
DB 922 TTCAATCGGGCGGGGGGAG 941

RESULT 30
US-10-027-632-248985
; Sequence 248985, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; TITLE OF INVENTION: Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
```

```
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 248985
; LENGTH: 1073
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-248985

Query Match      76.4%; Score 16.8; DB 14; Length 1073;
Best Local Similarity 90.0%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTCGATCGGGCGGGCGGAG 21
DB 922 TTCAATCGGGCGGGGGGAG 941

RESULT 31
US-10-027-632-248986
; Sequence 248986, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; TITLE OF INVENTION: Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 248986
; LENGTH: 1073
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-248986

Query Match      76.4%; Score 16.8; DB 14; Length 1073;
Best Local Similarity 90.0%; Pred. No. 1.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTCGATCGGGCGGGCGGAG 21
DB 922 TTCAATCGGGCGGGGGGAG 941

RESULT 32
US-10-274-095-47
; Sequence 47, Application US/10274095
; Publication No. US20030120433A1
; GENERAL INFORMATION:
; APPLICANT: Yokota, Hiroki
; APPLICANT: Sun, Hui Bin
; TITLE OF INVENTION: Methods for Predicting Transcription
; TITLE OF INVENTION: Levels
```

```

; FILE REFERENCE: ARTI_0137US
; CURRENT APPLICATION NUMBER: US/10/274,095
; PRIOR FILING DATE: 2002-10-17
; PRIOR APPLICATION NUMBER: 60/329,961
; PRIOR FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 47
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: fragment
US-10-274-095-47

Query Match          74.5%; Score 16.4; DB 15; Length 18;
Best Local Similarity 94.4%; Pred. No. 1.5e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGGCGAGC 22
    |||||
Db 1 GATCGGGCGGGCGGCGATC 18
    |||||

RESULT 33
US-10-204-708-68
; Sequence 68, Application US/10204708
; Publication No. US20030141852A1
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPNEROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication
; TITLE OF INVENTION: by Assessing DNA Methylation
; FILE REFERENCE: 5013.1012
; CURRENT APPLICATION NUMBER: US/10/204,708
; CURRENT FILING DATE: 2003-05-06
; PRIOR APPLICATION NUMBER: PCT/EP01/03971
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: DE 10019058.8
; PRIOR FILING DATE: 2000-04-06
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 98
; SEQ ID NO 68
; LENGTH: 6368
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-204-708-68

Query Match          74.5%; Score 16.4; DB 13; Length 6368;
Best Local Similarity 94.4%; Pred. No. 1.5e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGGCGAGC 22
    |||||
Db 4207 GATCGGGCGGGCGGCGC 4224
    |||||

RESULT 34
US-10-311-455-1912
; Sequence 1912, Application US/10311455
; Publication No. US20030143606A1
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPNEROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication
; TITLE OF INVENTION: cytosine methylation
; FILE REFERENCE: 5013.1014
; CURRENT APPLICATION NUMBER: US/10/311,455
; CURRENT FILING DATE: 2002-12-16
; PRIOR APPLICATION NUMBER: PCT/EP01/07537
; PRIOR FILING DATE: 2001-07-02
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 2424
; SEQ ID NO 68
; LENGTH: 6368
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-311-455-2414

Query Match          74.5%; Score 16.4; DB 13; Length 6988;
Best Local Similarity 94.4%; Pred. No. 1.5e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAG 21
    |||||
Db 1297 CGAGCGGGCGGGCGGCGAG 1314
    |||||

RESULT 35
US-10-311-455-2414
; Sequence 2414, Application US/10311455
; Publication No. US20030143606A1
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPNEROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with the Immune System by Det
; TITLE OF INVENTION: cytosine methylation
; FILE REFERENCE: 5013.1014
; CURRENT APPLICATION NUMBER: US/10/311,455
; CURRENT FILING DATE: 2002-12-16
; PRIOR APPLICATION NUMBER: PCT/EP01/07537
; PRIOR FILING DATE: 2001-07-02
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 2424
; SEQ ID NO 2414
; LENGTH: 6988
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-311-455-2414

Query Match          74.5%; Score 16.4; DB 13; Length 6988;
Best Local Similarity 94.4%; Pred. No. 1.5e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAG 21
    |||||
Db 1297 CGAGCGGGCGGGCGGCGAG 1314
    |||||

RESULT 36
US-10-067-977-3
; Sequence 3, Application US/10067977
; Publication No. US20030157679A1
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPNEROCK, Christian
; APPLICANT: YAN, Chunhua and KE, Zhaoxi
; TITLE OF INVENTION: ISOLATED HUMAN KINASE PROTEINS, NUCLEIC

```



```
; TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN KINASE PROTEINS, AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CLO01313
; CURRENT APPLICATION NUMBER: US/10/067,977
; CURRENT FILING DATE: 2002-02-08
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 10573
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-067-977-3
Query Match 74.5%; Score 16.4; DB 13; Length 10573;
Best Local Similarity 94.4%; Pred. No. 1.4e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGAG-21
DB 2862 CGGTCGGGGCGGGCGAG 2879

RESULT 37
US-10-292-198-1
; Sequence 1, Application US/10292198
; Publication No. US20030157654A1
; GENERAL INFORMATION:
; APPLICANT: SHEN, Ben
; APPLICANT: LIU, Wen
; TITLE OF INVENTION: BIOSYNTHESIS OF ENEDIYNE COMPOUNDS BY MANIPULATION OF C-1027 GENE
; FILE REFERENCE: 054030-0007
; CURRENT APPLICATION NUMBER: US/10/292,198
; CURRENT FILING DATE: 2003-03-14
; PRIOR APPLICATION NUMBER: US 10/159,257
; PRIOR FILING DATE: 2002-05-31
; PRIOR APPLICATION NUMBER: US 09/478,188
; PRIOR FILING DATE: 2000-01-05
; PRIOR APPLICATION NUMBER: US 60/115,434
; PRIOR FILING DATE: 1999-01-06
; NUMBER OF SEQ ID NOS: 146
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 1
; LENGTH: 63158
; TYPE: DNA
; ORGANISM: Streptomyces globisporus
US-10-292-198-1
Query Match 74.5%; Score 16.4; DB 13; Length 63158;
Best Local Similarity 94.4%; Pred. No. 1e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 GATCGGGCGGGCGAGC 22
DB 46469 GAGCGGGCGGGCGAGC 46486

RESULT 38
US-09-918-995-11259/c
; Sequence 11259, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FROM VARIOUS CDNA LIBRARIES
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 11259
```

```
; LENGTH: 498
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(498)
; OTHER INFORMATION: n = A,T,C or G
US-09-918-995-11259
```

```
Query Match 73.6%; Score 16.2; DB 11; Length 498;
Best Local Similarity 85.7%; Pred. No. 2.7e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
```

```
QY 2 TTCGATCGGGCGGGCGGAGC 22
DB 26 TTCTATTGGGGCGGGCGTGC 6
```

```
RESULT 39
US-10-156-761-1713
; Sequence 1713, Application US/10156761
; Publication No. US20030119018A1
; GENERAL INFORMATION:
; APPLICANT: OMURA, SATOSHI
; APPLICANT: IKEDA, HARUO
; APPLICANT: ISHIKAWA, JUN
; APPLICANT: HORIKAWA, HIROSHI
; APPLICANT: SHIBA, TADAYOSHI
; APPLICANT: SAKAKI, YOSHIYUKI
; APPLICANT: HATTORI, MASAHIRA
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-262
; CURRENT APPLICATION NUMBER: US/10/156,761
; CURRENT FILING DATE: 2002-05-29
; PRIOR APPLICATION NUMBER: JP 2001-204089
; PRIOR FILING DATE: 2001-05-30
; PRIOR APPLICATION NUMBER: JP 2001-272697
; PRIOR FILING DATE: 2001-08-02
; NUMBER OF SEQ ID NOS: 15109
; SEQ ID NO 1713
; LENGTH: 1599
; TYPE: DNA
; ORGANISM: Streptomyces avermitilis
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1599)
US-10-156-761-1713
```

```
Query Match 73.6%; Score 16.2; DB 15; Length 1599;
Best Local Similarity 85.7%; Pred. No. 2.3e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
```

```
QY 2 TTCGATCGGGCGGGCGGAGC 22
DB 607 TTCGAACGGGGCGGTGGCGGC 627
```

```
RESULT 40
US-09-769-207A-1
; Sequence 1, Application US/09769207A
; Patent No. US20020132234A1
; GENERAL INFORMATION:
; APPLICANT: DZGenes, LLC
; TITLE OF INVENTION: NITRIC OXIDE SYNTHASE GENE DIAGNOSTIC POLYMORPHISMS
; FILE REFERENCE: DZG 2165.1
; CURRENT APPLICATION NUMBER: US/09/769,207A
; CURRENT FILING DATE: 2001-01-24
; PRIOR APPLICATION NUMBER: US 60/177,775
; PRIOR FILING DATE: 2000-01-24
; PRIOR APPLICATION NUMBER: US 60/220,662
; PRIOR FILING DATE: 2000-07-25
; NUMBER OF SEQ ID NOS: 25
; SOFTWARE: PatentIn version 3.0
```

```

; SEQ ID NO 1
; LENGTH: 3586
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1465)..(3585)
; OTHER INFORMATION: Promotor region and exon 1, partial CDS
; NAME/KEY: mRNA
; LOCATION: (3473)..(3585)
; NAME/KEY: exon
; LOCATION: (3473)..(3586)
; NAME/KEY: Gene
; LOCATION: (3473)..(3585)
; OTHER INFORMATION: gene=NOS3
; NAME/KEY: CDS
; LOCATION: (3494)..(3586)
; OTHER INFORMATION: n=unknown
; US-09-769-207A-1

```

```

Query Match          73.6%; Score 16.2; DB 10; Length 3586;
Best Local Similarity 85.7%; Pred.No. 2e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

```

```

Qy      1 ATTCGATCGGGGGGGGGGAG 21
Db      3360 ATGGGATAGGGGGGGGGGAG 3380

```

```

Search completed: February 18, 2004, 17:16:29
Job time : 201 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 18, 2004, 16:10:46 ; Search time 1633 Seconds
(without alignments)
327.433 Million cell updates/sec

Title: US-10-026-341A-2

Perfect score: 22

Sequence: 1 attgatcggggcggggcggc 22

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST.*

1: em_estba.*

2: em_esthum.*

3: em_estin.*

4: em_estmu.*

5: em_estov.*

6: em_estpl.*

7: em_estro.*

8: em_hic.*

9: gb_est1.*

10: gb_est2.*

11: gb_hic.*

12: gb_est3.*

13: gb_est4.*

14: gb_est5.*

15: em_estfun.*

16: em_estom.*

17: em_gss_hum.*

18: em_gss_hiv.*

19: em_gss_pln.*

20: em_gss_vrt.*

21: em_gss_fun.*

22: em_gss_mam.*

23: em_gss_mus.*

24: em_gss_pro.*

25: em_gss_rod.*

26: em_gss_phg.*

27: em_gss_vrl.*

28: gb_gss1.*

29: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | ID | Description |
|------------|-------|-------------|--------|----|-------------|
| C 1 | 20 | 90.9 | 429 | 9 | AT005447 |
| 2 | 19 | 86.4 | 311 | 10 | B5654158 |
| C 3 | 18.4 | 83.6 | 1640 | 12 | BM805163 |
| 4 | 17.8 | 80.9 | 490 | 28 | BH629785 |

| | | | | | |
|------|------|------|------|----|----------|
| C 5 | 17.8 | 80.9 | 495 | 14 | CA708962 |
| C 6 | 17.8 | 80.9 | 552 | 29 | CC042414 |
| C 7 | 17.8 | 80.9 | 603 | 14 | CD228564 |
| C 8 | 17.8 | 80.9 | 710 | 29 | CD228564 |
| 9 | 17.8 | 80.9 | 723 | 29 | BZ997350 |
| C 10 | 17.8 | 80.9 | 896 | 14 | BZ339055 |
| 11 | 17.8 | 80.9 | 1511 | 12 | BZ339055 |
| C 12 | 17.8 | 80.9 | 1557 | 11 | BM552949 |
| 13 | 17.4 | 79.1 | 319 | 14 | AY106281 |
| 14 | 17.4 | 79.1 | 322 | 14 | CD550345 |
| 15 | 17.4 | 79.1 | 350 | 13 | CD550569 |
| 16 | 17.4 | 79.1 | 350 | 13 | BY302682 |
| 17 | 17.4 | 79.1 | 355 | 13 | BY210660 |
| C 18 | 17.4 | 79.1 | 375 | 9 | AY122220 |
| C 19 | 17.4 | 79.1 | 379 | 9 | AT846527 |
| 20 | 17.4 | 79.1 | 406 | 9 | AA615769 |
| 21 | 17.4 | 79.1 | 418 | 14 | CB765932 |
| 22 | 17.4 | 79.1 | 423 | 12 | BI220387 |
| 23 | 17.4 | 79.1 | 437 | 10 | BB859799 |
| 24 | 17.4 | 79.1 | 448 | 9 | AI194235 |
| C 25 | 17.4 | 79.1 | 472 | 14 | CB640552 |
| 26 | 17.4 | 79.1 | 493 | 10 | BE650594 |
| 27 | 17.4 | 79.1 | 523 | 4 | BX529827 |
| 28 | 17.4 | 79.1 | 530 | 10 | BF015640 |
| 29 | 17.4 | 79.1 | 600 | 13 | BU919508 |
| 30 | 17.4 | 79.1 | 604 | 13 | BU924221 |
| 31 | 17.4 | 79.1 | 624 | 14 | CB578330 |
| 32 | 17.4 | 79.1 | 635 | 14 | BY721913 |
| 33 | 17.4 | 79.1 | 645 | 14 | BY721911 |
| 34 | 17.4 | 79.1 | 679 | 12 | BI11370 |
| 35 | 17.4 | 79.1 | 704 | 29 | BZ658953 |
| 36 | 17.4 | 79.1 | 709 | 13 | BQ445767 |
| 37 | 17.4 | 79.1 | 735 | 13 | BQ444735 |
| 38 | 17.4 | 79.1 | 741 | 13 | BQ445216 |
| 39 | 17.4 | 79.1 | 764 | 12 | BI146925 |
| 40 | 17.4 | 79.1 | 773 | 14 | BY712338 |
| 41 | 17.4 | 79.1 | 774 | 10 | BG172820 |
| 42 | 17.4 | 79.1 | 776 | 13 | BQ746273 |
| C 43 | 17.4 | 79.1 | 781 | 14 | CB657329 |
| 44 | 17.4 | 79.1 | 791 | 12 | BI100531 |
| 45 | 17.4 | 79.1 | 793 | 13 | BUS25924 |

ALIGNMENTS

| | | | | | |
|------------|---|-------------|---------------------|----------------------|-----------------|
| RESULT 1 | AT005447/c | 429 bp | mRNA | linear | EST 25-MAR-2002 |
| LOCUS | AT005447 | POMF01 | Pleurotus ostreatus | cdna clone MFB34-F01 | mrna |
| DEFINITION | AT005447 | sequence. | | | |
| ACCESSION | AT005447 | GI:13420306 | | | |
| VERSION | AT005447 | EST. | | | |
| KEYWORDS | EST. | | | | |
| SOURCE | AT005447.1 | | | | |
| ORGANISM | Pleurotus ostreatus (oyster mushroom) | | | | |
| REFERENCE | Pleurotus ostreatus | | | | |
| AUTHORS | Eukaryota; Fungi; Basidiomycota; Hymenomycetes; Homobasidiomycetes; Agaricales; Pleurotaceae; Pleurotus. | | | | |
| TITLE | 1 (bases 1 to 429) | | | | |
| JOURNAL | Lee, S.H., Kim, B.G., Kim, K.J., Lee, J.S., Yun, D.W., Hahn, J.H., Kim, G.H., Lee, K.H., Suh, D.S., Kwon, S.T., Lee, C.S. and Yoo, Y.B. | | | | |
| MEDLINE | Comparative Analysis of Sequences Expressed during the Liquid-Cultured Mycelia and Fruit Body Stages of Pleurotus | | | | |
| PUBMED | 11849675 | | | | |
| COMMENT | Fungal Genet. Biol. 35 (2), 115-134 (2002) | | | | |

Email: bgkimyes@da.go.kr
Submitted through BRIC(Biological Research Information Center) of Korea

URL: http://bric.postech.ac.kr/
GeneNuri No. KS105130.

FEATURES

source
Location/Qualifiers

1. 429
/organism="Pleurotus ostreatus"
/mol_type="rRNA"
/cultivar="ASI 2029"
/db_xref="taxon:5322"
/clone="MFB34-F01"
/dev_stage="mature fruiting body"
/lab_host="E.coli"
/clone_lib="POMFBO1"
/note="Vector: lambda TriPEX2; Site 1: SfiI; Site 2: SfiI; average insert size:1500 bp; initial pu:5' 10 7; isolation of total RNA from the mature fruiting body cultivated in poplar tree sawdust bottle"

BASE COUNT 101 a 135 c 92 g 101 t

ORIGIN

Query Match 90.9%; Score 20; DB 9; Length 429;
Best Local Similarity 100.0%; Pred. No. 2.9e+02;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TTCGATCGGGGGGGCGAG 21

Db 120 TTCGATCGGGGGGGCGAG 101

RESULT 2

BE654158

LOCUS

DEFINITION BE654158 311 bp mRNA linear EST 06-SEP-2000
UI-M-ANI-aff-f-04-0-UI-r2 NIH BMAP MEG_N Mus musculus cDNA clone

ACCESSION BE654158

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

Genome Res. 6 (9), 791-806 (1996)

889548

Contact: Chin, H

National Institute of Mental Health

6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD

20892-9643, USA

Tel: 301 443 1706

Fax: 301 443 9890

Email: m8est@mail.nih.gov

CDNA Library Preparation: M.B. Soares Lab Clone distribution:

Researchers may obtain BMAP cDNA clones from RESEARCH GENETICS. It should be noted that Bento Soares is generating a small number of additional specialized non-redundant arrays of BMAP cDNAs whose availability will be considered under appropriate and limited collaborative arrangements. The following repetitive elements were found in this cDNA sequence: 119-174, >(CA)n#simple_repeat

Seq primer: M13 Reverse.

Location/Qualifiers

1. 311

/organism="Mus musculus"

/mol_type="rRNA"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UI-M-ANI-aff-f-04-0-UI"

/dev_stage="27-32 days"

FEATURES

source

BASE COUNT 222 a 508 c 438 g 369 t

ORIGIN

Query Match 83.6%; Score 18.4; DB 12; Length 1640;
Best Local Similarity 95.0%; Pred. No. 1e+03;

/lab_host="DH10B (Life Technologies)"

/clone_lib="NIH BMAP MEG_N"

/note="Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; The NIH BMAP MEG_N library is a normalized library constructed from mouse basal ganglia. The tag is a string of 5 nucleotides present between the Not I site and the oligo-dT track. The library was constructed as described by Bonaldo, Lennon and Soares, Genome Research 6: 791-806, 1996. Tissue provided by Ms. Annie Novakovich, Zivic-Miller Laboratories."

BASE COUNT 62 a 62 c 116 g 71 t

ORIGIN

Query Match 86.4%; Score 19; DB 10; Length 311;

Best Local Similarity 100.0%; Pred. No. 7.5e+02;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 CGATCGGGGGGGGGCGAG 22

Db 217 CGATCGGGGGGGGGCGAG 235

RESULT 3

BM805163/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BM805163

AGENCOURT 6490256 NIH_MGC_125 Homo sapiens cDNA clone IMAGE:5726046

5', mRNA sequence.

BM805163

BM805163.1 GI:19121986

EST.

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BM805163

AGENCOURT 6490256 NIH_MGC_125 Homo sapiens cDNA clone IMAGE:5726046

5', mRNA sequence.

BM805163

BM805163.1 GI:19121986

EST.

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BM805163

AGENCOURT 6490256 NIH_MGC_125 Homo sapiens cDNA clone IMAGE:5726046

5', mRNA sequence.

BM805163

AGENCOURT 6490256 NIH_MGC_125 Homo sapiens cDNA clone IMAGE:5726046

5', mRNA sequence.

BM805163

AGENCOURT 6490256 NIH_MGC_125 Homo sapiens cDNA clone IMAGE:5726046

5', mRNA sequence.

BM805163

AGENCOURT 6490256 NIH_MGC_125 Homo sapiens cDNA clone IMAGE:5726046

5', mRNA sequence.

BM805163

AGENCOURT 6490256 NIH_MGC_125 Homo sapiens cDNA clone IMAGE:5726046

5', mRNA sequence.

```

Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGCA 20
Db 1456 ATTCGATCGGGCGGGCGCA 1437

RESULT 4
BH629785
LOCUS
DEFINITION 490 bp DNA linear GSS 30-JAN-2002
1007082B12.2EL_Y1 1007 - RescueMu Grid H Zea mays genomic, genomic
survey sequence.
ACCESSION BH629785
VERSION BH629785
KEYWORDS GSS.
SOURCE BH629785.1 GI:18443036
ORGANISM Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE 1 (bases 1 to 490)
AUTHORS Walbot,V.
JOURNAL Maize genomic sequences found using engineered RescueMu transposon
COMMENT Unpublished
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Possible ligation site of ends cut by 2 different endonucleases.
Reverse complemented post-ligation sequence from source sequence.
Plate: 1007082 column: 31
Class: transposon-tagged.
FEATURES
source
Location/Qualifiers
1..490
/organism="Zea mays"
/mol_type="genomic DNA"
/cultivar="mixed background W23/A188/B73"
/db_xref="taxon:4577"
/tissue_type="leaf"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="1007 - RescueMu Grid H"
/note="Organ: Leaf; Vector: RescueMu (engineered from
pBluescript backbone); Site 1: BamHI; Site 2: BglII;
RescueMu is a 4.9 kb, modified maize Mu transposon
designed to allow plasmid rescue from total genomic DNA.
Mu elements insert preferentially into transcription
units. For more information on RescueMu, go to the web
site 'www.zmndb.iastate.edu' and follow the links for
'RescueMu.' Grid H was grown at Berkeley in 2001. DNA
was extracted from leaf punches, double digested using
BamHI and BglII, and ligated to form circular plasmids.
DH10B cells were transformed and then screened on LB
plates with ampicillin."
BASE COUNT 76 a 153 c 196 g 65 t
ORIGIN
Query Match 80.9%; Score 17.8; DB 28; Length 490;
Best Local Similarity 90.5%; Pred. No. 2.1e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 TTTCGATCGGGCGGGCGGAGC 22
Db 345 TTTCGATCGGGCGGGCGGAGC 365

RESULT 5
CA708962/c
LOCUS
DEFINITION 495 bp mRNA linear EST 26-NOV-2002
wdk2c.pk011.a20 wdk2c Triticum aestivum cDNA clone wdk2c.pk011.a20

```

```

5' end, mRNA sequence.
CA708962
CA708962.1, GI:25430755
EST.
Triticum aestivum (bread wheat)
Triticum aestivum
Triticum aestivum
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
; Triticeae; Triticum.
1 (bases 1 to 495)
Tingey,S.V., Powell,W., Wolters,P., Dolan,M., Hainey,C., Yuan,Z.,
Miao,G., Caraher,N. and Hanafey,M.X.
DuPont Wheat cDNA Sequence
Unpublished
Contact: Scott V. Tingey
Crop Genetics
E. I. DuPont de Nemours and Company
1 Innovation Way, P.O. Box 6104, Newark, DE 19714-6104, USA
Tel: 302-631-2602
Fax: 302-631-2607
Email: Scott.V.Tingey@USA.dupont.com
Seq primer: M13.
Location/Qualifiers
1..495
/organism="Triticum aestivum"
/mol_type="mRNA"
/db_xref="taxon:4565"
/clone="wdk2c.pk011.a20"
/tissue_type="kernel"
/clone_lib="wdk2c"
/note="Vector: pBluescript SK+; Site_1: EcoRI; Site_2:
XhoI; Wheat (Triticum aestivum L.) developing kernel, 7
days after anthesis."
BASE COUNT 91 a 165 c 165 g 71 t 3 others
ORIGIN
Query Match 80.9%; Score 17.8; DB 14; Length 495;
Best Local Similarity 90.5%; Pred. No. 2.1e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 ATTCGATCGGGCGGGCGGAG 21
Db 76 ATTCGATCGGGCGGGCGGAG 56

RESULT 6
CC042414/c
LOCUS
DEFINITION 3591_1_151_1_D02.Y.1 3591 - RescueMu Grid P Zea mays genomic,
genomic survey sequence.
ACCESSION CC042414
VERSION CC042414.1 GI:29457305
KEYWORDS GSS.
SOURCE CC042414.1
ORGANISM Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
1 (bases 1 to 552)
Walbot,V.
Maize genomic sequences found using engineered RescueMu transposon
Unpublished
JOURNAL
Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 2227
Fax: 650 725 8221
Email: walbot@stanford.edu
Plate: 3591_1_151_1 row: 28
Class: transposon-tagged.
Location/Qualifiers
1..552

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BZ339055
LOCUS 2339055 723 bp DNA linear GSS 06-NOV-2003
DEFINITION ic29d10.g1 WGS-Sbicolorf (JM107 adapted methyl filtered) Sorghum
bicolor genomic clone ic29d10 5', genomic survey sequence.
ACCESSION BZ339055
VERSION BZ339055.1 GI:24735457
KEYWORDS GSS.
SOURCE Sorghum bicolor (sorghum)
ORGANISM Sorghum bicolor
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Sorghum.
REFERENCE 1 (bases 1 to 723)
AUTHORS Rabinowicz,P.D., O'Shaughnessy,A.L., Balija,V., Dedhia,N.,
Katzenburger,F., King,L., Miller,B., Moller,S., Nascimento,L.,
Zutavern,T., Palmer,L., McCombie,W.R. and Martienssen,R.A.
Genomic shotgun sequences from Sorghum bicolor (methyl-filtered)
Unpublished
Contact: W. Richard McCombie
Lita Annenberg Hazen Genome Sequencing Center
Cold Spring Harbor Laboratory
PO Box 100, Cold Spring Harbor, NY 11724, USA
Tel: 516 367 8884
Fax: 516 367 8874
Email: mcombie@cshl.org
Plate: ic29 row: d column: 10
Seq primer: -21M13UnivRev
Class: shotgun
High quality sequence stop: 723.
Location/Qualifiers
1..723
/organism="Sorghum bicolor"
/mol_type="genomic DNA"
/db_xref="taxon:4558"
/clone="ic29d10"
/lab_host="JM107 or DH5a"
/clone_lib="WGS-Sbicolorf (JM107 adapted methyl filtered)"
/notes="Site 1: Xba I; Site 2: Xba I; The vector was
digested with XbaI and one nucleotide was added by fill in
in the recessive 3' end. The genomic DNA was rebluized,
end repaired, adaptor ligated and size fractionated using
sephadex. The resulting fragments were between 0.8 and 3
kb and were cloned into the vector (x/y reads in M13mp19,
.b/g reads in pUC19). The same ligation was transformed in
either JM107 or DH5a."
BASE COUNT 153 a 189 c 217 g 163 t 1 others
ORIGIN
Query Match 80.9%; Score 17.8; DB 29; Length 723;
Best Local Similarity 90.5%; Pred.No.2e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
CY 2 TTCGATCGGGCGGGCGGCGAGC 22
|||||
Db 78 TTCGATCGGAGCGGCGGCGAGC 98
|||||
RESULT 10
CD439046/c
LOCUS CD439046 896 bp mRNA linear EST 03-JUN-2003
DEFINITION EL01N0520C10.b EndospERM_5 Zea mays cDNA, mRNA sequence.
ACCESSION CD439046
VERSION CD439046.1 GI:31354689
KEYWORDS EST.
SOURCE Zea mays
ORGANISM Zea mays
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
clade; Panicoideae; Andropogoneae; Zea.
REFERENCE 1 (bases 1 to 896)
AUTHORS Lai,J., Dey,N., Kim,C.S., Becraft,P., Larkins,B., Linton,E. and
Messing,J.
Sequencing of the maize endospERM ESTs
TITLE

0.9-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 013. Note: this is a NIH MGC Library."

BASE COUNT 428 a 413 c 450 g 215 t 5 others

ORIGIN

Query Match 80.9%; Score 17.8; DB 12; Length 1511;
Best Local Similarity 90.5%; Pred. No. 1.7e+03;
Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 TTCGATCGGGCGGGCGGAGC 22

Db 54 TTCGACCGGGCGGGCGGAGC 74

RESULT 12

AY106281/c 1557 bp mRNA linear HTC 16-OCT-2002
LOCUS AY106281 PC0117884 mRNA sequence.

DEFINITION Zea mays

ACCESSION AY106281

VERSION AY106281.1 GI:21209359

KEYWORDS HTC

SOURCE Zea mays

ORGANISM Zea mays

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.

REFERENCE 1 (bases 1 to 1557)

AUTHORS Hainey, C.F., Dolan, M., Miao, G.H., Vogel, J.M., Whittitt, M.S., Arthur, L.W., Hanafey, M., Morgante, M. and Tingey, S.V.

TITLE Maize Mapping Project/DuPont Consensus Sequences for Design of Overgo Probes

JOURNAL Unpublished (2002)

REFERENCE 2 (bases 1 to 1557)

AUTHORS Coe, E.H.

TITLE Direct Submission

JOURNAL Submitted (25-APR-2002) Maize Mapping Project, University of Missouri, Columbia, MO 65211, USA

COMMENT If you are interested in getting corresponding physical clones, these are publicly available from ZmDB, www.zmdb.iastate.edu, TIGR, searching at MSL, maizemap.org; ZmDB, www.zmdb.iastate.edu; TIGR, www.tigr.org; or NCBI, www.ncbi.nlm.nih.gov. When the source of the maize cDNA sequences is either Virginia Walbot, Stanford or Pat Schnable, Iowa State, then clones may be requested from ZmDB: www.zmdb.iastate.edu.

FEATURES

source

Location/Qualifiers

1..1557

/organism="Zea mays"

/mol_type="mRNA"

/db_xref="MaizeDB:636873"

/db_xref="taxon:4577"

/clone_lib="Maize Mapping Project/DuPont Consensus Library"

/note="this sequence is part of a project of EST assemblies resulting from the application of public contigs to seed DuPont contigs; this resource was assembled by DuPont as part of a collaboration for the overgo addressing of BACs in conjunction with the Maize Mapping Project"

BASE COUNT 424 a 382 c 377 g 374 t

ORIGIN

Query Match 80.9%; Score 17.8; DB 11; Length 1557;

Best Local Similarity 90.5%; Pred. No. 1.7e+03;

Matches 19; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 TTCGATCGGGCGGGCGGAGC 22

Db 231 TTCGATCGGAGCGGGCGGAGC 211

RESULT 13

CD550345

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

FEATURES

source

Location/Qualifiers

1..319

/organism="Mus musculus"

/mol_type="mRNA"

/strain="C57BL/6J"

/db_xref="nlaEST:B0310F12-5"

/db_xref="taxon:10090"

/clone="NIA:B0310F12 IMAGE:30430055"

/tissue_type="E9.5 whole embryo"

/dev_stages="whole embryo including extraembryonic tissues at 9.5-days postcoitum"

/lab_host="DH10B"

/clone_lib="NIA Mouse E9.5 Whole Embryo cDNA Library (long)"

/note="Vector: pCMV-SPORT6 (Invitrogen); Site 1: SalI; Site 2: NotI; Mouse cDNA project by the Laboratory of Genetics, National Institute on Aging (NIA), Intramural Research Program, NIH (<http://igsun.grc.nia.nih.gov/cDNA>). This is a long-transcript enriched cDNA library (Ref. Genome Res. 11: 1553-1558 (2001). [PMID: 11544199]). Total RNAs were extracted from a pool of 16 embryos at 9.5-days postcoitum. Double-stranded cDNAs were synthesized with an oligo(dT) primer (Invitrogen): 5'-pGATCTAGTCGAGCGGCGGCGCTTTT-3' from 6.1 ug of total RNA, treated with T4 DNA polymerase, and purified by ethanol-precipitation. The cDNAs were ligated to lone-linker LL-Sal4, purified by phenol/chloroform, and separated from free linkers by Centricon 100. Then, the cDNAs were amplified by long-range high fidelity PCR using Ex Taq polymerase (Takara) with a primer Sal4-S. The products were purified by phenol/chloroform and Centricon 100. The cDNAs were digested with SalI and NotI enzymes and cloned into SalI/NotI site of pCMV-SPORT6 plasmid vector. The DH10B E. coli host was transformed with the ligation mixture by the standard chemical method. The average insert size is about 3.0Kb. The library was constructed by Yulan Piao."

BASE COUNT 73 a 87 c 108 g 51 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 14; Length 319;

Best Local Similarity 94.7%; Pred. No. 3.2e+03;

Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGAGC 22

CD550345 319 bp mRNA linear EST 11-JUN-2003

B0310F12-5 NIA Mouse E9.5 Whole Embryo cDNA Library (Long) Mus

musculus cDNA clone NIA:B0310F12 IMAGE:30430055 5', mRNA sequence.

CD550345

CD550345.1 GI:31598076

EST.

Mus musculus (house mouse)

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathu; Muridae; Murinae; Mus.

1 (bases 1 to 319)

Piao, Y., Ko, N.T., Lim, M.K. and Ko, M.S.H.

Construction of long-transcript enriched cDNA libraries from

submicrogram amounts of total RNAs by a universal PCR amplification

method

Genome Res. 11 (9), 1553-1558 (2001)

21429098

11544199

Contact: Dawood B. Dudekula

Laboratory of Genetics

National Institute on Aging/National Institutes of Health

333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA

Email: cdna@igsun.grc.nia.nih.gov

Plate: B0310 row: F column: 12

Seq primer: M13 Reverse

High quality sequence stop: 319

POLYA=No.


```

Db          60 CGAGCGGGCGGGCGGAGC 78
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Query Match 79.1%; Score 17.4; DB 14; Length 322;
Best Local Similarity 94.7%; Pred. No. 3.2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

RESULT 14
CD550569      322 bp      mRNA      linear      EST 11-JUN-2003
LOCUS      B0315A09-5 NIA Mouse E9.5 Whole Embryo cDNA Library (Long) Mus
DEFINITION  musculus cDNA clone NIA:B0315A09 IMAGE:30430472 5', mRNA sequence.
ACCESSION  CD550569
VERSION     CD550569.1 GI:31598300
KEYWORDS
SOURCE      Mus musculus (house mouse)
ORGANISM    Mus musculus

REFERENCE
AUTHORS     Piao,Y., Ko,N.T., Lim,M.K. and Ko,M.S.H.
TITLE       Construction of long-transcript enriched cDNA libraries from
            submicrogram amounts of total RNAs by a universal PCR amplification
            method
JOURNAL     Genome Res. 11 (9), 1553-1558 (2001)
MEDLINE     21429098
PUBMED      11544199
COMMENT     Contact: Dawood B. Dudekula
            Laboratory of Genetics
            National Institute on Aging/National Institutes of Health
            333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
            Email: cdna@nslu.grc.nia.nih.gov
            Plate: B0315 row: A column: 09
            Seq primer: M13 Reverse
            High quality sequence stop: 322
            POLYA=No.

FEATURES
            Location/Qualifiers
            1..322
            /organism="Mus musculus"
            /mol_type="mRNA"
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            /db_xref="nia:EST:B0315A09-5"
            /db_xref="taxon:10090"
            /clone="NIA:B0315A09 IMAGE:30430472"
            /tissue_type="E9.5 whole embryo"
            /dev_stage="whole embryo including extraembryonic tissues
            at 9.5-days postcoitum"
            /lab_host="DH10B"
            /clone_lib="NIA Mouse E9.5 Whole Embryo cDNA Library (Long
            )"
            /note="vector: pCMV-SPORT6 (Invitrogen); Site 1: SalI;
            Site 2: NotI; Mouse cDNA project by the Laboratory of
            Genetics, National Institute on Aging (NIA), Intramural
            Research Program, NIH (http://lgsun.grc.nia.nih.gov/cDNA).
            This is a long-transcript enriched cDNA library [Ref.
            Genome Res. 11: 1553-1558 (2001). [PMID: 11544199]]. Total
            RNAs were extracted from a pool of 16 embryos at 9.5-days
            postcoitum. Double-stranded cDNAs were synthesized with an
            oligo(dT) primer [Invitrogen:
            5'-pGACTAGTCAGATCGAGCGGCCGCCCTTTT-3']
            and from 6.1 ug of total RNA, treated with T4 DNA polymerase,
            and purified by ethanol-precipitation. The cDNAs were
            ligated to lone-linker linker 1L-Sal4, purified by
            phenol/chloroform, and separated from free linkers by
            Centricion 100. Then, the cDNAs were amplified by
            long-range high fidelity PCR using Ex Taq polymerase
            (Takara) with a primer Sal4-S. The products were purified
            by phenol/chloroform and Centricion 100. The cDNAs were
            digested with SalI and NotI enzymes and cloned into
            SalI/NotI site of pCMV-SPORT6 plasmid vector. The DH10B E.
            coli host was transformed with the ligation mixture by the
            standard chemical method. The average insert size is about
            3.0Kb. The library was constructed by Yulan Piao."
            74 a      88 c      109 g      51 t

BASE COUNT
ORIGIN

```

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4 CGATCGGGCGGGCGGAGC 22
|||||
Db          60 CGAGCGGGCGGGCGGAGC 78
|||||
Query Match 79.1%; Score 17.4; DB 14; Length 322;
Best Local Similarity 94.7%; Pred. No. 3.2e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

RESULT 15
BZ204565
LOCUS      BZ204565
DEFINITION  BZ204565 RIKEN full-length enriched, B6-derived CD11 +ve dendritic
            cells Mus musculus cDNA clone F730213P11 5', mRNA sequence.
ACCESSION  BZ204565
VERSION     BZ204565.1 GI:26384371
KEYWORDS
SOURCE      Mus musculus (house mouse)
ORGANISM    Mus musculus

REFERENCE
AUTHORS     Okazaki,Y., Furuno,M., Kasukawa,T., Adachi,J., Bono,H., Kondo,S.,
            Nikaido,I., Osato,N., Saito,R., Suzuki,H., Yamanaka,I., Kiyosawa,H.,
            Yagi,K., Tomaru,Y., Hasegawa,Y., Nogami,A., Schonbach,C.,
            Gocho,T., Baldarelli,R., Hill,D.P., Bult,C., Hume,D.A.,
            Quackenbush,J., Schriml,L.M., Kanapin,A., Matsuda,H., Batalov,S.,
            Beisel,K.W., Blake,J.A., Bradt,D., Bruscia,V., Chothia,C., Corbani
            L.E., Cousins,S., Dalla,E., Dragani,T.A., Fletcher,C.F., Forrest
            A., Frazer,K.S., Gaasterland,T., Gariboldi,M., Gissi,C., Godzik,A.,
            Gough,J., Grimmond,S., Gustincich,S., Hirokawa,N., Jackson,I.J.,
            Jarvis,E.D., Kanai,A., Kawaji,H., Kawasawa,Y., Kedzierski,R.M.,
            King,B.L., Konagaya,A., Kurochkin,I.V., Lee,Y., Lennard,B., Lyons
            P.A., Maglott,D.R., Maltais,L., Marchionni,L., McKenzie,L., Miki
            H., Nagashima,T., Numata,K., Okido,T., Pavan,W.J., Perceat,G.,
            Pesole,G., Petrovsky,N., Pillai,R., Pontius,J.U., Qi,D.,
            Ramachandran,S., Ravasi,T., Reed,J.C., Reed,D.J., Reid,J., Ring
            B.Z., Ringwald,M., Sandelin,A., Schneider,C., Semple,C.A., Secou
            M., Shimada,K., Sultana,R., Takenaka,Y., Taylor,M.S., Teasdale
            R.D., Tomita,M., Verardo,R., Wagner,L., Wahlestedt,C., Wang,Y.,
            Watanabe,Y., Wells,C., Wilming,L.G., Wynshaw-Boris,A., Yanagisawa
            M., Yang,I., Yang,L., Yuan,Z., Zavolan,M., Zhu,Y., Zimmer,A.,
            Carninci,P., Hayatsu,N., Hirozane-Kishikawa,T., Konno,H., Nakamura
            M., Sakazume,N., Sato,K., Shiraki,T., Waki,K., Kawai,J., Aizawa,K.,
            Arakawa,T., Fukuda,S., Hara,A., Hashizume,W., Imotani,K., Ishii
            Y., Itoh,M., Kagawa,I., Miyazaki,A., Sakai,K., Sasaki,D., Shibata
            K., Shingawa,A., Yasunishi,A., Yoshino,M., Waterston,R., Lander
            E.S., Rogers,J., Birney,E. and Hayashizaki,Y.
            Analysis of the mouse transcriptome based on functional annotation
            of 60,770 full-length cDNAs
            Nature 420, 563-573 (2002)
            22354683
            12466851
            Contact: Yoshihide Hayashizaki
            Laboratory for Genome Exploration Research Group, RIKEN Genomic
            Sciences Center (GSC), Yokohama Institute
            The Institute of Physical and Chemical Research (RIKEN)
            1-7-22 Sukeno-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
            Tel: 81-45-503-9222
            Fax: 81-45-503-9216
            Email: genome-res@gs.c.riken.go.jp/
            URL: http://genome.gsc.riken.go.jp/
            Aizawa,K., Akimura,T., Arakawa,T., Carninci,P., Fukuda,S., Hirozane
            T., Imotani,K., Ishii,Y., Itoh,M., Kawai,J., Konno,H., Miyazaki,A.,
            Murata,M., Nakamura,M., Nomura,K., Numazaki,R., Ohno,M., Sakai,K.,
            Sakazume,N., Sasaki,D., Sato,K., Shibata,K., Shiraki,T., Tagami
            M., Waki,K., Watahiki,A., Muramatsu,M. and Hayashizaki,Y. Direct
            Submission
            Computational Analysis of Full-length Mouse cDNAs Compared with
            Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
            Normalization and subtraction of cap-trapper-selected cDNAs to
            prepare full-length cDNA libraries for rapid discovery of new
            genes. Genome Res. 10 (10), 1617-1630 (2000)

```

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Tissues were provided by Dr. John Todd (Dept. of Medical Genetics Wellcome Trust Centre for Molecular Mechanisms in Disease Wellcome Trust/MRC building Addenbrookes Hospital Cambridge) whose assistance we gratefully acknowledge.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

FEATURES

source

Location/Qualifiers

1. 341
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="F730213P11"
/cell_type="B6-derived CD11 +ve dendritic cells"
/clone_lib="RIKEN full-length enriched, B6-derived CD11 +ve dendritic cells"

64 a 77 C 130 G 70 t

BASE COUNT

ORIGIN

Query Match 79.1%; Score 17.4; DB 13; Length 341;
Best Local Similarity 94.7%; Pred. No. 3.1e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY

4 CGATCGGGCGGGCGGCG 22

Db

136 CGAGCGGGCGGGCGGCG 154

RESULT 16

LOCUS

BY302682 350 bp mRNA linear EST 11-DEC-2002
R302682 RIKEN full-length enriched, 14.5 days embryo df/df
Rathke's pouches Mus musculus cDNA K82005D04 5', mRNA
sequence.

ACCESSION

BY302682.1 GI:26493019

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 350)
Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Oatso, N., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H.,
Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C.,
Gojobori, T., Baldarelli, R., Hill, B. P., Bult, C., Hume, D. C.,
Quackenbush, J., Schram, L. M., Kanapin, A., Matsuda, H., Batalov, S.,
Beisel, K. W., Blake, J. A., Bradt, D., Brusic, V., Chothia, C., Corbani,
L. E., Cousins, S., Dalla, E., Dragani, T. A., Fletcher, C. F., Forrest,
A., Frazer, K. S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A.,
Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I. J.,
Jarvis, E. D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R. M.,
King, B. L., Kongaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons,
P. A., Maglott, D. R., Maltais, L., Marchionni, L., McKenrie, J., Maki,
H., Nagashima, T., Numata, K., Okido, T., Pavan, W. J., Pertea, G.,
Pesole, G., Petrovsky, N., Pillai, R., Pontius, J. U., Qi, D.,
Ramachandran, S., Ravasi, T., Reed, J. C., Reed, D. J., Reid, J., Ring,
B. Z., Ringwald, M., Sanderlin, A., Schneider, C., Semple, C. A., Setou,
M., Shmada, K., Sultana, R., Takenaka, Y., Taylor, M. S., Teasdale,
R. D., Tomita, M., Verardo, R., Wagner, L., Wahlstedt, C., Wang, Y.,
Watanabe, Y., Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yang, L.,
M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A.,

Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K., Arawaka, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shingawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E. S., Rogers, J., Birney, E. and Hayashizaki, Y.

Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs

Nature 420, 563-573 (2002)

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

12466851
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216

Email: genome-res@gsc.riken.go.jpURL: <http://genome.gsc.riken.go.jp/>

Aizawa, K., Akimura, T., Arawaka, T., Carninci, P., Fukuda, S., Hirozane, T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Konno, H., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Waki, K., Watabiki, A., Muramatsu, M. and Hayashizaki, Y. Direct

Submission

Computational Analysis of Full-length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Tissues were provided by Michelle Brinkmeier and Sally Camper (Dept. Human Genetics University of Michigan Medical School 4301 MSRB 3 1500 W. Medical Center Dr. Ann Arbor, MI 48109-0638 USA) whose assistance we gratefully acknowledge.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

FEATURES

source

Location/Qualifiers

1. 350
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="K82005D04"
/tissue_type="Rathke's pouches"
/dev_stage="14.5 days embryo df/df"
/clone_lib="RIKEN full-length enriched, 14.5 days embryo
df/df Rathke's pouches"

BASE COUNT 77 a 84 C 127 G 62 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 13; Length 350;
Best Local Similarity 94.7%; Pred. No. 3.1e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCG 22

Db 111 CGAGCGGGCGGGCGGCG 129

RESULT 17

BY210660

LOCUS

BY210660

355 bp

mRNA

linear

EST 10-DEC-2002

Wellcome Trust Centre for Molecular Mechanisms in Disease Wellcome
Trust/MRC building Addenbrookes Hospital Cambridge) whose
assistance we gratefully acknowledge.
Please visit our web site (<http://genome.gsc.riken.go.jp>) for
further details.

| | | |
|-----------------------|--|--|
| FEATURES | source | Location/Qualifiers |
| | | 1. 355 |
| | | /organism="Mus musculus" |
| | | /mol_type="mRNA" |
| | | /strain="C57BL/6J" |
| | | /db_xref="taxon:10090" |
| | | /clone="F730317H16" |
| | | /cell_type="B6-derived CD11 +ve dendritic cells" |
| | | /clone_lib="RIKEN full-length enriched, B6-derived CD11 +ve dendritic cells" |
| BASE COUNT | 70 a 79 c 133 g 71 t 2 others | |
| ORIGIN | | |
| Query Match | 79.1%; Score 17.4; DB 13; Length 355; | |
| Best Local Similarity | 94.7%; Pred. No. 3.le-03; | |
| Matches | 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0; | |
| QY | 4 CGATCGGGGGCGGGCGAGC 22 | |
| | | |
| DB | 136 CGAGCGGGGGCGGGCGAGC 154 | |
| | | |
| RESULT 18 | | |
| LOCUS | AW122220/c | |
| DEFINITION | AW122220 375 bp mRNA linear EST 22-OCT-1999 | |
| VERSION | UI-M-BH2.2-av-d-07-0-UI.s1 NIH BMAP M.S3.2 Mus musculus cDNA clone | |
| KEYWORDS | UI-M-BH2.2-av-d-07-0-UI 3', mRNA sequence. | |
| SOURCE | AW122220 GI:6037683 | |
| ORGANISM | EST. | |
| | Mus musculus (house mouse) | |
| REFERENCE | | |
| AUTHORS | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. | |
| TITLE | 1 (bases 1 to 375) | |
| | Bonaldo, M.F., Lennon, G. and Soares, M.B. | |
| | Normalization and subtraction: two approaches to facilitate gene | |
| | discovery | |
| JOURNAL | Genome Res. 6 (9), 791-806 (1996) | |
| MEDLINE | 97044477 | |
| PUBMED | 8899548 | |
| COMMENT | Contact: Chin, H | |
| | National Institute of Mental Health | |
| | 6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD | |
| | 20892-9643, USA | |
| | Tel: 301 443 1706 | |
| | Fax: 301 443 9890 | |
| | Email: mstet@mail.nih.gov | |
| | Oligo-dT track not found, Not I site shown in beginning of sequence | |
| | is likely internal to the message. cDNA Library Preparation: M.B. | |
| | Soares Lab Clone Distribution: NIH BMAP cDNA clones will be made | |
| | available by the means that is soon to be determined. When NIH | |
| | determines the means for distribution of the BMAP cDNA clones, this | |
| | record will be updated accordingly when that means is determined. | |
| | The following repetitive elements were found in this cDNA sequence | |
| | 192-247, >(CA)n#simple repeat | |
| | Seq primer: M13 Forward | |
| | POLYA=No. | |
| FEATURES | Location/Qualifiers | |
| | 1. 375 | |
| | /organism="Mus musculus" | |
| | /mol_type="mRNA" | |
| | /strain="C57BL/6J" | |
| | /db_xref="taxon:10090" | |
| | /clone="UI-M-BH2.2-av-d-07-0-UI" | |
| | /dev_stages="27-32 days" | |
| | /lab_host="DH10B (Life Technologies)" | |
| | /clone_lib="NIH BMAP M.S3.2" | |

| | | |
|--|--|---|
| | BV210660 | RIKEN full-length enriched, B6-derived CD11 +ve dendritic cells Mus musculus cDNA clone F73031H16 5', mRNA sequence. |
| DEFINITION | | |
| ACCESSION | BV210660 | |
| VERSION | | |
| KEYWORDS | | |
| SOURCE | BV210660.1 GI:26391233 | |
| ORGANISM | Mus musculus (house mouse) | |
| Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; | | |
| Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. | | |
| 1 (bases 1 to 355) | | |
| OKazaki,Y., Furuno,M.N., Kasukawa,T., Adachi,J., Bono,H., Kondo,S., Nikaido,I., Osato,N., Saito,R., Suzuki,H., Yamanaka,T., Kiyosawa,H., Yagi,K., Tomaru,Y., Hasegawa,Y., Nogami,A., Schonbach,C., Gojobori,T., Baldarelli,L.M., Kanpin,A., Matsuda,H., Batalov,S., Quackenbush,J., Schriml,R.H., Bratt,D.J., Brusci,V., Chochia,C., Corbi Beisel,K.W., Blake,J.A., Bradt,D.V., Fletcher,C.F., Forrest, L.E., Cousins,S., Dalla,E.S., Dragani,T.A., Goss,C., Godrik,A. A., Frazer,K.S., Gaasterland,T., Gariboldi,M., Glass,C., Godzik,A. Gough,J., Grimmond,S., Gustincich,S., Hirokawa,N., Jackson,I.J., Jarvis,E.D., Kanai,A., Kawaji,H., Kawasaki,Y., Kedzierski,R.M., King,B.L., Konggay,A., Kurachkin,I.V., Lee,Y., Lenhard,B., Lyons P.A., Maglott,D.R., Maltais,L., Marchionni,L., McKenzie,L., Miki H., Nagashima,T., Numata,K., Okido,T., Pavan,W.J., Perle,G., Pesole,G., Petrovsky,N., Pillai,R., Pontius,J.U., Reid,J., Ring Ramchandran,S., Ravasi,T., Reed,J.C., Reed,D.J., Seid,J., RamaChandran,M., Sandelin,A., Schneider,C., Semple,C.A., Secou M., Shimada,K., Sultan,R., Takenaka,Y., Taylor,M.S., Teasdale R.D., Tonita,M., Verdaro,R., Wagner,L., Wahlestedt,C., Wang,Y., Watanabe,Y., Wells,C., Wilming,L.G., Wynshaw-Boris,A., Yanagisawa M., Yang,I., Yang,L., Yuan,Z., Zavolan,M., Zhu,Y., Zimmer,A., Ishii M., Sakazume,N., Sato,K., Shiraki,T., Waki,K., Kwai,J., Aizawa,K., Arakawa,T., Fukuda,S., Harai,A., Hashizume,W., Imotani,K., Ishii K., Itoh,M., Kagawa,I., Miyazaki,A., Sakai,K., Sasaki,D., Shibata Y., Shingawa,A., Yasunishi,A., Yoshino,M., Waterston,R., Lander E.S., Rogers,J., Birney,B. and Hayashizaki,Y. | | |
| TITLE | | Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs |
| JOURNAL | Nature 420, 563-573 (2002) | |
| MEDLINE | 22354683 | |
| PUBMED | 12466881 | |
| COMMENT | Contact: Yoshihide Hayashizaki Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute The Institute of Physical and Chemical Research (RIKEN) 1-7-2 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222 Fax: 81-45-503-9216 Email: genome-res@gsc.riken.go.jp/ URL:http://genome.gsc.riken.go.jp/ Aizawa,K., Akimura,T., Arakawa,T., Carninci,P., Fukuda,S., Hirozane T., Imotani,K., Ishii,Y., Itoh,M., Kawai,J., Konno,H., Miyazaki,A. Murata,M., Nakamura,M., Nomura,K., Numazaki,R., Ohno,M., Sekai, Sakazume,N., Sakai,D., Sato,K., Shibata,K., Shiraki,T., Tagami M., Waki,K., Watabiki,A., Muramatsu,M. and Hayashizaki,Y. Direct Submission Computational Analysis of Full-length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001) Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes Genome Res. 10 (10) 1617-1630 (2000) RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000) Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001) cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. Tissues were provided by Dr. John Todd (Dept. of Medical Genetics | |

/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; The NIH-BMAP M.S3.2 library is a subtracted library of a series, ultimately derived from a mixture of individually tagged normalized libraries from ten regions of the mouse brain (cerebellum, brain stems, olfactory bulbs, hypothalamus, cortex, amygdala, basal ganglia, pineal gland, striatum, hippocampus) after a series of subtractions to reduce the representation of cDNAs from which ESTs had already been generated. The following serially subtracted libraries were generated in this process: NIH-BMAP M.S3.2, NIH-BMAP M.S2, NIH-BMAP M.S1. The subtracted library (NIH-BMAP M.S3.2) was constructed as follows: PCR-amplified cDNA inserts from NIH-BMAP M.S2 clones from which 3' ESTs had been derived was used as a driver in a hybridization with the NIH-BMAP M.S2 library in the form of single-stranded circles. The remaining single-stranded circles (subtracted library) was purified by hydroxyapatite column chromatography, converted to double-stranded circles and electroporated into DH10B bacteria (Life Technologies) to generate the NIH-BMAP M.S3.2 library. This procedure has been previously described (Bonaldo, Lennon and Soares, Genome Research 6:791-806, 1996)

Research 6:791-806, 1996)
TAG LIB=NIH-BMAP M.S3.2
TAG-TISSUE=Cerebellum
TAG-SEQ=GACTC"

BASE COUNT 87 a 136 c 77 g 75 t
ORIGIN

Query Match 79.1%; Score 17.4; DB 9; Length 375;
Best Local Similarity 94.7%; Pred. No. 3.1e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGAGC 22
Db 100 CGAGCGGGCGGGCGGAGC 82

RESULT 19
LOCUS A1846527/c
DEFINITION UI-M-ANI-aff-f-04-0-UI.s1 NIH-BMAP M.S3.2 Mus musculus cDNA clone
UI-M-ANI-aff-f-04-0-UI 3', mRNA sequence.

ACCESSION A1846527.1 GI:5490433
VERSION A1846527
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 379)
AUTHORS Bonaldo,M.F., Lennon,G. and Soares,M.B.
TITLE Normalization and subtraction: two approaches to facilitate gene discovery

JOURNAL Genome Res. 6 (9), 791-806 (1996)
MEDLINE 97044477
PubMed 8889548

COMMENT Contact: Chin, H
National Institute of Mental Health
6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD 20892-9643, USA
Tel: 301 443 1706
Fax: 301 443 9890
Email: m3st@mail.nih.gov
Oligo-dt track not found, Not I site shown in beginning of sequence is likely internal to the message. cDNA library preparation: M.B. Soares Lab Clone distribution: NIH-BMAP cDNA clones will be made available by the means that is soon to be determined. When NIH determines the means for distribution of the BMAP cDNA clones, this record will be updated accordingly when that means is determined. The following repetitive elements were found in this cDNA sequence: 192-247, >(CA)n\$Simple_repeat

Seq primer: M13 Forward
POLYA=No.

FEATURES
source

1..379
/organism="Mus musculus"
/mol_type="mRNA"
/strain="CS7BL/6J"
/db_xref="taxon:10090"
/clone="UI-M-ANI-aff-f-04-0-UI"
/dev_stage="27-32 days"
/lab_host="DH10B (Life Technologies)"
/clone_lib="NIH-BMAP M.S3.2"
/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; The NIH-BMAP M.S3.2 library is a normalized library constructed from mouse basal ganglia. The tag is a string of 5 nucleotides present between the Not I site and the oligo-dt track. The library was constructed as described by Bonaldo, Lennon and Soares, Genome Research 6: 791-806, 1996. Tissue provided by Ms. Annie Novakovich, Zivic-Miller Laboratories.
TAG LIB=NIH-BMAP M.S3.2
TAG-TISSUE=Hippocampus
TAG-SEQ=ITCGA"

BASE COUNT 89 a 136 c 77 g 77 t
ORIGIN

Query Match 79.1%; Score 17.4; DB 9; Length 379;
Best Local Similarity 94.7%; Pred. No. 3.1e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGGGCGGGCGGAGC 22
Db 100 CGAGCGGGCGGGCGGAGC 82

RESULT 20
LOCUS AA615769
DEFINITION v072f08.r1 Barstead mouse myctubus MPRB5 Mus musculus cDNA clone
IMAGE:1064679 5', similar to WP:C16C10.7 CE01498 ZINC FINGER PROTEIN
/, mRNA sequence.

ACCESSION AA615769
VERSION AA615769.1 GI:2502997
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 406)
AUTHORS Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T., Geisler,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M., Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B., Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and Waterston,R.
TITLE The WashU-HMI Mouse EST Project
JOURNAL Unpublished
COMMENT Contact: Marra M/Mouse EST Project
WashU-HMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@wustl.edu
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
MG1:587039
Seq primer: -28m13 rev2 ET from Amersham
High quality sequence stop: 398.
Location/Qualifiers
1..406
/organism="Mus musculus"
/mol_type="mRNA"

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| | BEST | linear | EST 11-JUL-2001 |
| | B1220387 | 423 bp mRNA | |
| | LOCUS | 602935686F1 NCI_CGAP_Li9 Mus musculus cDNA clone IMAGE:5098916 5' | |
| | DEFINITION | mRNA sequence. | |
| | ACCESSION | B1220387 | |
| | VERSION | B1220387.1 GI:14673931 | |
| | KEYWORDS | EST. | |
| | SOURCE | Mus musculus (house mouse) | |
| | ORGANISM | Mus musculus | |
| | REFERENCE | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus; | |
| AUTHORS | NIH-MGC | http://mgc.nci.nih.gov/ National Institutes of Health, Mammalian Gene Collection (MGC) Unpublished Contact: Robert Strauszberg, Ph.D. Email: cgabbs+email.nih.gov | |
| TITLE | CNA Library Preparation: | Jeffrey E. Green, M.D. | |
| JOURNAL | CNA Library Arrayed by: | The I.M.A.G.E. Consortium (LLNL) | |
| COMMENT | DNA Sequencing by: | Incyte Genomics, Inc. Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Plate: LLAMil1237 row: 1 column: 21 High quality sequence stop: 419. | |

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/organism="Mus musculus"
/mol_type="mRNA"
/_strain="FVB/N"
/db_xref="taxon:10090"
/_clone="IMAGE:5098516"
/_lab_host="DH10B (T1 phage-resistant)"
/_clone_lib="NCI_CGAP_L19"
/notes="Organ: liver; Vector: pCW-SPORT6; Site_1: NotI; Site_2: SalI; Cloned unidirectionally. Primer: Oligo dt. Average insert size 1.9 kb. Constructed by Life Technologies. Note: this is a NCI_CGAP Library."

BASE COUNT      88 a   104 c   155 g      76 t
ORIGIN
Query Match      79.1%;   Score 17.4;   DB 12;   Length 423;
Best Local Similarity 94.7%;   Pred. No.3e+03;
Matches 18;   Conservative 0;   Mismatches 1;   Indels 0;   Gaps 0;

QY      4   CGATCGGGGGCGGGGCGGAGC 22
      ||| ||||| ||||| |||||
DB     138  CGAGCGGGGGCGGGGCGGAGC 156

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| | |
|------------|---|
| RESULT 23 | |
| BB859799 | |
| LOCUS | 437 bp mRNA linear EST 26-NOV-2001 |
| DEFINITION | BB859799 RIKEN full-length enriched, kidney CCH-142 RAG CDNA YMA musculus cdna clone G43001A01 5', mRNA sequence. |
| ACCESSION | BB859799 |
| VERSION | BB859799.1 GI:17101253 |
| KEYWORDS | EST. |
| SOURCE | Mus musculus (house mouse) |
| ORGANISM | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. |
| REFERENCE | 1 (bases 1 to 437) |
| AUTHORS | Akimura,T., Arakawa,T., Carninci,P., Furuno,M., Hanagaki,T., Hayatsu,N., Hiramoto,K., Hizaoka,T., Hirozane,T., Imotani,K., Ishii,Y., Ito,M., Kawai,J., Kojima,Y., Konno,H., Kouda,M., Matsuyama,T., Nakamura,M., Nishi,K., Nomura,K., Numasaki,R., Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sakazume,N., Sasaki,D., Sato,K., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagawa, |

TITLE RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al. 2001)

JOURNAL COMMENT Unpublished
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsc.riken.go.jp,
URL: http://genome.gsc.riken.go.jp/
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M., and Hayashizaki, Y., et al.
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)
wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuurra S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--394-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)
Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y., and Hayashizaki, Y.
Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
Please visit our web site (http://genome.gsc.riken.go.jp) for further details.
e mouse tissues.

FEATURES Location/Qualifiers
1..437
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="G43001CA01"
/tissue_type="kidney"
/cell_line="CCL-142 RAG"
/clone_lib="RIKEN full-length enriched, kidney CCL-142 RAG cDNA"

BASE COUNT 89 a 103 c 160 g 85 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 10; Length 437;
Best Local Similarity 94.7%; Pred. No. 3e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
|||||
Db 168 CGAGCGGGCGGGCGGAGC 186
|||||

RESULT 24
A1194235
LOCUS 448 bp mRNA linear EST 13-OCT-1998
DEFINITION ue52ell.r1 Soares mammary_gland NMLMG Mus musculus cDNA clone IMAGE:1494764 5', similar to TR:035445 O35445 HYPOTHETICAL 19.8 KD PROTEIN. ;, mRNA sequence.

ACCESSION A1194235
VERSION A1194235.1 GI:3745442
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 448)
Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T., Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M., Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B., Theisinger, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and Waterston, R.

TITLE The WashU-HMI Mouse EST Project

JOURNAL COMMENT Unpublished
Contact: Marra M/Mouse EST Project
WashU-HMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@wustl.edu
This clone is available royalty-free through LLNL ; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:932368
Seq primer: -28ml3 rev2 ET from Amersham.

FEATURES Location/Qualifiers
1..448
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="IMAGE:1494764"
/sex="female (lactating)"
/tissue_type="mammary gland"
/lab_host="DH10B"
/clone_lib="Soares mammary_gland NMLMG"
/note="Vector: pTV3D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from mammary gland tissue from a lactating female, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pTV3 vector. Library is normalized. Library was constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 90 a 110 c 161 g 87 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 9; Length 448;
Best Local Similarity 94.7%; Pred. No. 3e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
|||||
Db 143 CGAGCGGGCGGGCGGAGC 161
|||||

RESULT 25
CB640552/c
LOCUS 472 bp mRNA linear EST 08-APR-2003
DEFINITION OSJNEa15E16.f OSJNEA Oryza sativa (japonica cultivar-group) cDNA clone OSJNEa15E16 5', mRNA sequence.

ACCESSION CB640552
VERSION CB640552.1 GI:29635543
KEYWORDS EST.
SOURCE Oryza sativa (japonica cultivar-group)
ORGANISM Oryza sativa (japonica cultivar-group)
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Ehrhartoideae; Oryzeae; Oryza.

REFERENCE 1 (bases 1 to 472)
Jantasuriyarat, C., Lu, G., Gowda, M., Hatfield, J., Zhou, B., Mazur, E., Kudrna, D., Dean, R., Soderlund, C., Wing, R. and Wang, G.
Large-scale identification of ESTs involved in the interaction between rice and Magnaporthe grisea

JOURNAL COMMENT Unpublished
Contact: Rod Wing
Arizona Genomics Institute
University of Arizona
Biological Sciences West, 448A, P.O. Box 210088, Tucson, AZ 85721-0088, USA
Tel: 520 626 3967
Fax: 520 621 9288
Email: http://genome.arizona.edu
PCR Primers
FORWARD: atc agc ggc cgc gat cc
BACKWARD: aat taa ccc tca cta aag gg

```

Plate: 15 row: E column: 16
Seq primer: atc agc ggc cgc gat cc.
Location/Qualifiers
1. .472
/organism="Oryza sativa (japonica cultivar-group)"
/mol_type="RNA"
/cultivar="Nipponbare"
/db_xref="taxon:39947"
/clone="OSUNEAL5E16"
/dev_stage="3 week"
/lab_host="DH10B"
/clone_lib="OSNEA"
/note="Vector: pBluescript II KS +; Site 1: EcoRI; Site 2:
XhoI; 6 hrs after immunoculation with Rice Blast (Che 86061
)"
BASE COUNT      79 a 146 c 166 g 81 t
ORIGIN
Query Match      79.1%; Score 17.4; DB 14; Length 472;
Best Local Similarity 94.7%; Pred. No. 3e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 TCGATCGGGCGGGCGGAG 21
|||||
Db 43 TCGATCGGGCGGGCGGAG 25

RESULT 26
BE650594
LOCUS
DEFINITION
UI-M-BH2.2-aov-d-01-0-UI.r1 NIH BMAP M.S3.2 Mus musculus cDNA clone
UI-M-BH2.2-aov-d-01-0-UI 5', mRNA sequence.
ACCESSION
BE650594
VERSION
BE650594.1 GI:9976418
KEYWORDS
EST.
SOURCE
Mus musculus (house mouse)
ORGANISM
Mus musculus
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
REFERENCE
1 (bases 1 to 493)
Bonaldo,M.F., Lennon,G. and Soares,M.B.
Normalization and subtraction: two approaches to facilitate gene
discovery
Genome Res. 6 (9), 791-806 (1996)
9704477
MEDLINE
PUBMED
8889548
COMMENT
Contact: Chin, H
National Institute of Mental Health
6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD
20892-9643, USA
Tel: 301 443 1706
Fax: 301 443 9890
Email: mEST@mail.nih.gov
cDNA Library Preparation: M.B. Soares Lab Clone distribution:
Researchers may obtain BMAP cDNA clones from RESEARCH GENETICS. It
should be noted that Bento Soares is generating a small number of
additional specialized non-redundant arrays of BMAP cDNAs whose
availability will be considered under appropriate and limited
collaborative arrangements. The following repetitive elements were
found in this cDNA sequence: 118-173, >(CA)n#Simple_repeat
Seq primer: M13 Reverse.
Location/Qualifiers
1. .493
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UI-M-BH2.2-aov-d-01-0-UI"
/dev_stage="27-32 days"
/lab_host="DH10B (Life Technologies)"
/clone_lib="NIH BMAP M.S3.2"
/note="Vector: pT73D-Pac (Pharmacia) with a modified

```

polylinker; Site 1: Not I; Site 2: Eco RI; The NIH BMAP M.S3.2 library is a subtracted library of a series, ultimately derived from a mixture of individually tagged normalized libraries from ten regions of the mouse brain (cerebellum, brain stems, olfactory bulbs, hypothalamus, cortex, amygdala, basal ganglia, pineal gland, striatum, hippocampus) after a series of subtractions to reduce the representation of cDNAs from which ESTs had already been generated. The following serially subtracted libraries were generated in this process: NIH BMAP M.S2, NIH BMAP M.S2, NIH BMAP M.S1. The subtracted library (NIH BMAP M.S3.2) was constructed as follows: PCR amplified cDNA inserts from NIH BMAP M.S2 clones from which 3' ESTs had been derived was used as a driver in a hybridization with the NIH BMAP M.S2 library in the form of single-stranded circles. The remaining single-stranded circles (subtracted library) was purified by hydroxyapatite column chromatography, converted to double-stranded circles and electroporated into DH10B bacteria (Life Technologies) to generate the NIH BMAP M.S3.2 library. This procedure has been previously described (Bonaldo, Lennon and Soares, Genome Research 6:791-806, 1996)"

BASE COUNT 99 a 112 c 158 g 124 t
ORIGIN

Query Match 79.1%; Score 17.4; DB 10; Length 493;
Best Local Similarity 94.7%; Pred. No. 3e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGAGC 22
|||||
Db 265 CGAGCGGGCGGGCGGAGC 283

RESULT 27

BX529827
ID BX529827 standard; RNA; EST; 523 BP.

XX AC BX529827;

XX SV BX529827.1

XX DT 27-MAY-2003 (Rel. 75, Created)

XX DT 27-MAY-2003 (Rel. 75, Last updated, Version 1)

XX DE RZPD Mus musculus cDNA clone IMAGp952G154 = IMAGE:336469 5' EST.

XX EST; expressed sequence tag.

XX OS Mus musculus (house mouse)

XX OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia;

XX OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

XX RN [1]

XX RP 1-523

XX RA Hell O., Ebert L., Neubert P., Peters M., Radelof U., Schneider D.,

XX RA Korn B.;

XX RT Submitted (28-MAY-2003) to the EMBL/GenBank/DBJ databases.

XX RL RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH Im Neuenheimer

XX RL Feld 580, D-69120 Heidelberg, Germany

XX CC RZPD; IMAGp952G154.

XX CC RZPDLib; I.M.A.G.E. cDNA Clone Collection;

XX CC Mouse Unigeneset - RZPD2 (RZPDLib No.981)

XX CC http://www.rzpd.de/CloneCards/cgi-bin/showLib.pl.cgi/response?libNo=981

XX CC Contact: Ina Rolfs

XX CC RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH

XX CC Heubnerweg 6, D-14059 Berlin, Germany

XX CC Tel: +49 30 32639 101

XX CC Fax: +49 30 32639 111

XX CC www.rzpd.de

FEATURES
source

CC This clone is available royalty-free from RZPD;
 CC contact RZPD (clone@rzpd.de) for further information.
 CC Seq primer: T7, Primer sequence: TAATAGGACTCACTATAGG

XX Key Location/Qualifiers

FT source 1..523
 FT /db_xref="taxon:10090"
 FT /note="1st strand cDNA was primed with a Not I - oligo(dT)
 FT primer [5'
 FT TGTTCACCAATGCTGAAGTGGGAGCGCGCATGTCTTTTTTTTTTTT 3', on
 FT total mouse RNA [provided by Minoru Ko, Wayne State Univ.];
 FT double-stranded cDNA was ligated to EcoRI adaptors
 FT 5'-AATTCGGGACAGG-3' and 5'-CTCTGTCGG-3' (Pharmacia),
 FT digested with NotI and cloned into the NotI and EcoRI sites
 FT of the pT73D-Paci vector. Library went through one round
 FT of normalization, and was constructed by Bento Soares and
 FT M. Fatima Bonaldo."
 FT /organism="Mus musculus"
 FT /clone="IMAGp952G154"
 FT /clone_lib="Soares 19.5dpc p3NMF19.5"
 FT /dev_stage="19.5dpc total fetus"
 FT /lab_host="DH10B"

SQ Sequence 523 BP; 110 A; 126 C; 186 G; 101 T; 0 other;

Query Match 79.1%; Score 17.4; DB 4; Length 523;
 Best Local Similarity 94.7%; Pred. No. 2.9e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAGC 22

|||||

DB 237 CGAGCGGGCGGGCGGCGAGC 255

RESULT 28

BF015640

LOCUS

DEFINITION BF015640 530 bp mRNA linear EST 29-DEC-2000
 uy27b02.y1 NCI CGAP Lu30 Mus musculus cDNA clone IMAGE:3660747 5,
 similar to TR:O35445 O35445 HYPOTHETICAL 19.8 KD PROTEIN. ;, mRNA
 sequence.

ACCESSION BF015640.1 GI:10746972

VERSION

KEYWORDS

SOURCE

ORGANISM Mus musculus (house mouse)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

1 (bases 1 to 530)

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: Gilbert Smith, Ph.D.

cDNA Library Preparation: Life Technologies, Inc.

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

image.llnl.gov/image/html/resources.shtml

MGI:1421515

Seq primer: -40RP from Gibco

High quality sequence stop: 457.

Location/Qualifiers

source

1..530

/organism="Mus musculus"

/mol_type="mRNA"

/strain="CZECH II"

/db_xref="taxon:10090"

/clone="IMAGE:3660747"

/tissue_type="tumor, metastatic to mammary"
 /lab_host="DH10B"

/clone_lib="NCI-CGAP Lu30"

/note="Organ: lung; Vector: pCMV-SPORT6; Site: 1: NotI;
 Site 2: SalI; transgenic model WNT-1, expression driven by
 MMTV-LTR enhancer; Cloned unidirectionally. Primer: Oligo
 dT. Library constructed by Life Technologies.

Investigator providing samples: Gilbert Smith, NIH"

BASE COUNT 101 a 135 c 179 g 115 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 10; Length 530;

Best Local Similarity 94.7%; Pred. No. 2.9e+03;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAGC 22

|||||

DB 144 CGAGCGGGCGGGCGGCGAGC 162

RESULT 29

BU919508

LOCUS

DEFINITION BU919508 600 bp mRNA linear EST 30-OCT-2002
 6024-75 Mouse E14.5 retina lambda ZAP II Library Mus musculus cDNA,
 mRNA sequence.

ACCESSION BU919508

VERSION

KEYWORDS

SOURCE

ORGANISM

Mus musculus (house mouse)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 600)

Mu.X., Zhao, S., Perashad, R., Haieh, T.-F., Scarpa, A., Wang, S.W.,
 White, R.A., Beremand, P.D., Thomas, T.L., Gan, L. and Klein, W.H.

Gene expression in the developing mouse retina by EST sequencing
 and microarray analysis

Nucleic Acids Res. 29 (24), 4983-4993 (2001)

21671825

11812828

Contact: Klein WH

Department of Biochemistry and Molecular Biology

University of Texas M.D. Anderson Cancer Center

Box 117, 1515 Holcombe Blvd., Houston, TX 77030, USA

Tel: 713 792 3646

Fax: 713 790 0329.

Location/Qualifiers

1..600

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/tissue_type="neural retina"

/dev_stage="embryonic day 14.5 post-fertilization"

/clone_lib="Mouse E14.5 retina lambda ZAP II Library"

/clone 161 c 197 g 128 t

BASE COUNT 114 a 161 c 197 g 128 t

ORIGIN

Query Match 79.1%; Score 17.4; DB 13; Length 600;

Best Local Similarity 94.7%; Pred. No. 2.9e+03;

Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAGC 22

|||||

DB 134 CGAGCGGGCGGGCGGCGAGC 152

RESULT 30

BU924221

LOCUS

DEFINITION BU924221 600 bp mRNA linear EST 30-OCT-2002
 7082-67 Mouse E14.5 retina lambda ZAP II Library Mus musculus cDNA,
 mRNA sequence.

ACCESSION BU924221

VERSION

BU924221.1 GI:24428104


```

KEYWORDS
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 600)
Nu,X., Zhao,S., Pershad,R., Hsieh,T.-F., Scarpa,A., Wang,S.W.,
White,R.A., Beremand,P.D., Thomas,T.L., Gan,L. and Klein,W.H.
TITLE Gene expression in the developing mouse retina by EST sequencing
and microarray analysis
JOURNAL Nucleic Acids Res. 29 (24), 4983-4993 (2001)
MEDLINE 21671825
PUBMED 11812828
COMMENT Contact: Klein WH
Department of Biochemistry and Molecular Biology
University of Texas M.D. Anderson Cancer Center
Box 117, 1515 Holcombe Blvd., Houston, TX 77030, USA
Tel: 713 792 3646
Fax: 713 790 0329
FEATURES
source Location/Qualifiers
1..600
/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/tissue_type="neural retina"
/dev_stage="embryonic day 14.5 post-fertilization"
/clone_lib="Mouse E14.5 retina lambda ZAP II Library"
BASE COUNT 120 a 157 c 192 g 126 t 5 others
ORIGIN
Query Match 79.1%; Score 17.4; DB 13; Length 600;
Best Local Similarity 94.7%; Pred. No. 2.9e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 4 CGATCGGGCGGGCGGCGAGC 22
|||
DB 131 CGAGCGGGCGGGCGGCGAGC 149
|||

RESULT 31
CB578330 624 bp mRNA linear EST 03-APR-2003
LOCUS ANGNNUC:NRDGI-00162-E5-A nrdg1 (10855) Rattus norvegicus cDNA clone
DEFINITION nrdg1-00162-e5 5', mRNA sequence.
CB578330
CB578330.1 GI:29522371
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM Rattus norvegicus (Norway rat)
REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 624)
Angen EST Program.
Angen Rat EST Program
JOURNAL Unpublished
COMMENT Contact: Dan Fitzpatrick
Angen, Inc
One Angen Center Drive, Thousand Oaks, CA 91320-1799, USA
Tel: 805 447-4881
Plate: 00162 row: e column: 5.
FEATURES
source Location/Qualifiers
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/organism="Rattus norvegicus"
/mol_type="mRNA"
/db_xref="taxon:10116"
/clone_lib="nrdg1-00162-e5"
/tissue_type="Dorsal Root Ganglia"
/clone_lib="nrdg1 (10855)"
/notes="vector: pSPOR1; Site_1: SalI; Site_2: NotI; rat
dorsal root ganglia"
BASE COUNT 113 a 174 c 201 g 135 t 1 others
ORIGIN

```

```

Query Match 79.1%; Score 17.4; DB 14; Length 624;
Best Local Similarity 94.7%; Pred. No. 2.9e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 4 CGATCGGGCGGGCGGCGAGC 22
|||
DB 112 CGAGCGGGCGGGCGGCGAGC 130
|||

RESULT 32
BY721913 635 bp mRNA linear EST 17-DEC-2002
LOCUS BY721913
DEFINITION BY721913 RIKEN full-length enriched, 12 days embryo, embryonic body
between diaphragm region and neck Mus musculus cDNA clone
943090J12 5', mRNA sequence.
BY721913
BY721913.1 GI:27135030
EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 635)
Okazaki,Y., Furuno,M., Kasukawa,T., Adachi,J., Bono,H., Kondo,S.,
Nikaido,I., Omato,N., Saito,R., Suzuki,H., Yamataka,I., Kiyosawa,H.
Yagi,K., Tomaru,Y., Hasegawa,Y., Nogami,A., Schonbach,C.,
Gojobori,T., Baldarelli,R., Hill,D.P., Bult,C., Hume,D.A.,
Quackenbush,J., Schriml,L.M., Kanapin,A., Matsuda,H., Batalov,S.,
Beisel,K.W., Blake,J.A., Bradt,D., Brusic,V., Ciothia,C., Corbani
,L.E., Cousins,S., Dalla,E., Dragan,T.A., Fletcher,C.F., Forrest
,A., Frazer,K.S., Gaasterland,T., Gariboldi,M., Gissi,C., Godzik,A.
Gough,J., Grimmond,S., Gustincich,S., Hirokawa,N., Jackson,I.J.,
Jarvis,E.D., Kanai,A., Kawaji,H., Kawasawa,Y., Kedzierski,R.M.,
King,B.L., Konagaya,A., Kurochkin,I.V., Lee,Y., Lenhard,B., Lyons
,P.A., Maglott,D.R., Maltais,L., Marchionni,L., McKenzie,L., Miki
,H., Nagashima,T., Numata,K., Okido,T., Pavan,W.J., Pertea,G.,
Pesole,G., Petrovsky,N., Pillai,R., Pontius,D.J., Qi,D., Ring
Ramachandran,S., Ravasi,T., Reed,J.C., Reed,D.J., Reid,J., Ring
,B.Z., Ringwald,M., Sandelin,A., Schneider,C., Sempile,C.A., Setou
,M., Shingawa,K., Sultana,R., Takenaka,Y., Taylor,M.S., Teasdale
,R.D., Tomita,M., Verardo,R., Wagner,L., Wahlestedt,C., Wang,Y.,
Watanabe,Y., Wells,C., Wilming,L.G., Wyszewski,B., Zhu,Y., Zimmer
,M., Yang,Y., Yang,L., Yuan,Z., Zavolan,M., Zhu,Y., Zimmer
,M., Hayatsu,N., Hirozane-Kishikawa,T., Kono,H., Nakamura
,M., Sakazume,N., Sato,K., Shiraki,T., Waki,K., Kawai,J., Kawai,K.,
Arakawa,T., Fukuda,S., Hara,A., Hashizume,W., Imotani,K., Ishii
,Y., Itoh,M., Kagawa,I., Miyazaki,A., Sakai,K., Sasaki,D., Shibata
,K., Shinagawa,A., Yasunishi,A., Yoshino,M., Waterston,R., Lander
,E.S., Rogers,J., Birney,E. and Hayashizaki,Y.
ANALYSIS of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
JOURNAL
MEDLINE 22354683
PUBMED 12466851
COMMENT Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suhiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsc.riken.go.jp,
URL:http://genome.gsc.riken.go.jp/
ADACHI,J., Aizawa,K., Akimura,T., Arakawa,T., Carninci,P., Fukuda
,S., Hashizume,W., Hayashida,K., Hirozane,T., Hori,F., Imotani,K.,
Ishii,Y., Itoh,M., Kagawa,I., Kawai,J., Kawai,Y., Kondo,S., Konno
,H., Koya,S., Miyazaki,A., Murata,M., Nakamura,M., Nomura,K.,
Numazaki,R., Ohno,M., Ohsato,N., Saito,R., Sakazume,N., Sano,H.,
Sasaki,D., Sato,K., Shibata,K., Shiraki,T., Tagami,M., Takeda,Y.,
Waki,K., Watanuki,A., Muramatsu,M. and Hayashizaki,Y. Direct
Computational Analysis of Full-Length Mouse cDNAs Compared with

```

Gojobori, T., Baldarelli, R., Hill, D. P., Bult, C., Hume, D. A., Quackenbush, J., Schriml, L. M., Kanpin, A., Matsuda, H., Batalov, S., Baissel, K. W., Blake, J. A., Bradt, D., Brusic, V., Chochia, C., Corbani, L. E., Cousins, S., Dalla, E., Dragani, T. A., Fletcher, C. F., Forrest, A. A., Frazer, K. S., Gaasterland, T., Gariboldi, M., Glasi, C., Godzik, A., Gough, J., Grimmond, S., Gustinch, S., Hirokawa, N., Jackson, I. J., Jarvis, E. D., Kawaji, H., Kawasawa, Y., Kedzierski, R. M., King, B. L., Konagaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons, P. A., Maglott, D. R., Maltais, L., Marchionni, L., McKenzie, J., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W. J., Pertea, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J. U., Qi, D., Ramachandran, S., Ravasi, T., Reed, J. C., Reed, D. J., Reid, J., Ring, B. Z., Ringwald, M., Sadelain, A., Schneider, C., Semple, C. A., Setou, M., Shmada, K., Sultana, R., Takenaka, Y., Taylor, M. S., Teasdale, R. D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yanagisawa, M., Yang, I., Yang, L., Yuan, Z., Zavoian, M., Zhu, Y., Zimmerman, J., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Akizawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shinagawa, A., Yasunishi, A., Yoshino, M., Yatsushiro, R., Landier, E. S., Rogers, J., Birney, E. and Hayashizaki, Y.

Analysis of the mouse transcriptome based on functional annotation of 60, 770 full-length cDNAs

Nature 420, 563-573 (2002)

Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-resgsc.riken.go.jp,
URL: http://genome.gsc.riken.go.jp/
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kawai, J., Kojima, Y., Konno, S., Konno, H., Koya, S., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N., Saito, R., Sakazume, N., Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Takeda, Y., Waki, K., Watanabe, A., Yamada, M., Yamashita, Y., Hayashizaki, Y. Direct Submission

Computational Analysis of Full-length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in Riken Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

| Location/Qualifiers |
|---|
| 1. .645 |
| /organism="Mus musculus" |
| /mol_type="mRNA" |
| /db_xref="taxon:10090" |
| /clone="9430089102" |
| /tissue_type="embryonic body between diaphragm region and neck" |
| /dev_stage="12 days embryo" |
| /lab_host="DH108" |

```

FEATURES
source
high quality sequence: yes.
Location/Qualifiers
1. 679
/organism="Mus musculus"
/mol_type="mRNA"
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/db_xref="taxon:10090"
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/clone_lib="NCI CGAP Mam5"
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Site 2: NotI. Cloned unidirectionally. Primer: Oligo dT.
Library constructed by Life Technologies. Investigators
providing samples: Lothar Hennighausen/Robin Humphreys,
NIH"
136 a 190 c 216 g 146 t 1 others
BASE COUNT

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| DEFINITION | BQ445767 709 bp mRNA linear EST 29-MAY-2002 UI-M-ERO-bxm-g-18-0-UI.r2 NIH BMAP_ERO Mus musculus CDNA clone |
| ACCESSION | IMAGE:5710121.5', mRNA sequence. |
| VERSION | BQ445767 |
| KEYWORDS | BQ445767.1 GI:21248879 |
| SOURCE | EST. |
| ORGANISM | Mus musculus (house mouse) |
| | Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciuronomathi; Muridae; Mus; |

REFERENCE 1 (bases 1 to 709)
 AUTHORS NIH-MGC <http://mgs.nci.nih.gov/>
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-remail.nih.gov
 Tissue Procurement: Dr. James Lin, University of Iowa
 CDNA Library Preparation: Dr. M. Bento Soares, University of Iowa
 CDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
 This clone was contributed by the Brain Molecular Anatomy Project (BMAP)

Seq primer: pYX-5.
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 /tissue_type="whole brain"
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 /lab_host="DH10B (T1 phage resistant)"
 /clone_lib="NIH BMAP ERO"
 /note="Organ: brain; Vector: pYX-Asc; Site 1: EcoR I; Site 2: Not I; The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. Denatured mRNA was size fractionated on a 1% agarose gel. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was size selected according to mRNA size fraction, ligated with EcoR I adaptor, digested with Not I, and then cloned directionally into pYX-Asc vector. The library tag sequence located between the Not I site and the polyA tail is GGGCGGGA. This library was created for the University of Iowa Mouse Brain Molecular Anatomy Project (BMAP); 'Gene Discovery in the Developing Mouse Nervous System', supported by National Institutes of Mental Health (NIMH), Hemin Chin, Ph.D., program coordinator."

BASE COUNT 131 a 201 c 211 g 164 t 2 others
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 Best Local Similarity 94.7%; Pred. No. 2.8e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGGGGGGGCGGAGC 22 735 bp mRNA linear EST 29-MAY-2002
 DB 90 CGAGCGGGGGGGGGCGGAGC 108
 IMAGE:5710148 5', mRNA sequence.

RESULT 37
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 LOCUS
 DEFINITION
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
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 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 735)
 NIH-MGC <http://mgs.nci.nih.gov/>
 National Institutes of Health, Mammalian Gene Collection (MGC)
 Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-remail.nih.gov
 Tissue Procurement: Dr. James Lin, University of Iowa

REFERENCE 1 (bases 1 to 709)
 AUTHORS NIH-MGC <http://mgs.nci.nih.gov/>
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-remail.nih.gov
 Tissue Procurement: Dr. James Lin, University of Iowa

CDNA Library preparation: Dr. M. Bento Soares, University of Iowa
 CDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
 This clone was contributed by the Brain Molecular Anatomy Project (BMAP)

Seq primer: pYX-5.
 Location/Qualifiers
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 /db_xref="taxon:10090"
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 /dev_stage="embryo 15.5 dpc"
 /lab_host="DH10B (T1 phage resistant)"
 /clone_lib="NIH BMAP ERO"
 /note="Organ: brain; Vector: pYX-Asc; Site 1: EcoR I; Site 2: Not I; The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. Denatured mRNA was size fractionated on a 1% agarose gel. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was size selected according to mRNA size fraction, ligated with EcoR I adaptor, digested with Not I, and then cloned directionally into pYX-Asc vector. The library tag sequence located between the Not I site and the polyA tail is GGGCGGGA. This library was created for the University of Iowa Mouse Brain Molecular Anatomy Project (BMAP); 'Gene Discovery in the Developing Mouse Nervous System', supported by National Institutes of Mental Health (NIMH), Hemin Chin, Ph.D., program coordinator."

BASE COUNT 135 a 205 c 225 g 168 t 2 others
 ORIGIN
 Query Match 79.1%; Score 17.4; DB 13; Length 735;
 Best Local Similarity 94.7%; Pred. No. 2.8e+03;
 Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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 DB 90 CGAGCGGGGGGGGGCGGAGC 108
 IMAGE:5711200 5', mRNA sequence.

RESULT 38
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 DEFINITION
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
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 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 741)
 NIH-MGC <http://mgs.nci.nih.gov/>
 National Institutes of Health, Mammalian Gene Collection (MGC)
 Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-remail.nih.gov
 Tissue Procurement: Dr. James Lin, University of Iowa
 CDNA Library Preparation: Dr. M. Bento Soares, University of Iowa
 CDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
 This clone was contributed by the Brain Molecular Anatomy Project

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(BMAP)
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/clone="IMAGE:5711200"
/tissue_type="whole brain"
/dev_stage="embryo 15.5 dpc"
/lab_host="DH10B (T1 phage resistant)"
/clone_lib="NIH_BMAP_ERO"
/notes="Organ: brain; Vector: pYX-Asc; Site 1: EcoR I; Site 2: Not I; The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. Denatured mRNA was size fractionated on a 1% agarose gel. First strand cDNA synthesis was primed with an oligo-dr primer containing a Not I site. Double stranded cDNA was size selected according to mRNA size fraction, ligated with EcoR I adaptor, digested with Not I, and then cloned directionally into pYX-Asc vector. The library tag sequence located between the Not I site and the polyA tail, is GTCGCGGAA. This library was created for the University of Iowa Mouse Brain Molecular Anatomy Project (BMAP): 'Gene Discovery in the Developing Mouse Nervous System', supported by National Institutes of Mental Health (NIMH), Hemin Chin, Ph.D., program coordinator."
BASE COUNT 133 a 202 c 224 g 179 t 3 others
ORIGIN
Query Match 79.1%; Score 17.4; DB 13; Length 741;
Best Local Similarity 94.7%; Pred. No. 2.8e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAGC 22
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DB 129 CGAGCGGGCGGGCGGCGAGC 147

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LOCUS 602511551F1 NCI_CGAP_Li9 Mus musculus cDNA clone IMAGE:5052660 5',
DEFINITION mRNA sequence.
ACCESSION BI146925
VERSION BI146925.1 GI:14606926
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 764)
Okazaki, F., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Otsu, R., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H.,
Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C.,
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L. E., Cousins, S., Dalla, E., Dragani, T. A., Fletcher, C. F., Forrest,
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Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I. J.,
Jarvis, E. D., Kanai, A., Kawai, H., Kawasawa, Y., Kedzierski, R. M.,
King, B. L., Konagaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons,
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R. D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y.,
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M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A.,
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M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K.,
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K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander,
E. S., Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
NATURE 420, 563-573 (2002)
22354683
12466851
PUBLISHED
COMMENT
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsr.riken.go.jp/
URL: http://genome.gsc.riken.go.jp/
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda

FEATURES
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Seq primer: pYX-5.
Location/Qualifiers
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/clone_lib="NIH_BMAP_ERO"
/notes="Organ: brain; Vector: pYX-Asc; Site 1: EcoR I; Site 2: Not I; The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. Denatured mRNA was size fractionated on a 1% agarose gel. First strand cDNA synthesis was primed with an oligo-dr primer containing a Not I site. Double stranded cDNA was size selected according to mRNA size fraction, ligated with EcoR I adaptor, digested with Not I, and then cloned directionally into pYX-Asc vector. The library tag sequence located between the Not I site and the polyA tail, is GTCGCGGAA. This library was created for the University of Iowa Mouse Brain Molecular Anatomy Project (BMAP): 'Gene Discovery in the Developing Mouse Nervous System', supported by National Institutes of Mental Health (NIMH), Hemin Chin, Ph.D., program coordinator."
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Query Match 79.1%; Score 17.4; DB 13; Length 741;
Best Local Similarity 94.7%; Pred. No. 2.8e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CGATCGGGCGGGCGGCGAGC 22
|||||
DB 129 CGAGCGGGCGGGCGGCGAGC 147

RESULT 39
BI146925 764 bp mRNA linear EST 05-JUL-2001
LOCUS 602511551F1 NCI_CGAP_Li9 Mus musculus cDNA clone IMAGE:5052660 5',
DEFINITION mRNA sequence.
ACCESSION BI146925
VERSION BI146925.1 GI:14606926
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 764)
Okazaki, F., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Otsu, R., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H.,
Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C.,
Gojobori, T., Baldarelli, R., Hill, D. P., Bult, C., Hume, D. A.,
Quackenbush, J., Schriml, L. M., Kanapin, A., Matsuda, H., Batalov, S.,
Beisel, K. W., Blake, J. A., Bradt, D., Brusci, V., Chothia, C., Corbani,
L. E., Cousins, S., Dalla, E., Dragani, T. A., Fletcher, C. F., Forrest,
A., Frazer, K. S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A.,
Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I. J.,
Jarvis, E. D., Kanai, A., Kawai, H., Kawasawa, Y., Kedzierski, R. M.,
King, B. L., Konagaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons,
P. A., Maglott, D. R., Maltais, L., Marchionni, L., McKenzie, L., Miki,
H., Nagashima, T., Numata, K., Okido, T., Pavan, W. J., Pertea, G.,
Pesole, G., Petrovsky, N., Pillai, R., Pontius, J. U., Qi, D.,
Ramachandran, S., Ravasi, T., Reed, J. C., Reed, D. J., Reid, J., Ring,
B. Z., Ringwald, M., Sandelin, A., Schneider, C., Sempile, C. A., Setou,
M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M. S., Teasdale,
R. D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y.,
Watanabe, Y., Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yanagisawa,
M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A.,
Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura,
M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K.,
Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii,
Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata,
K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander,
E. S., Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
NATURE 420, 563-573 (2002)
22354683
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PUBLISHED
COMMENT
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsr.riken.go.jp/
URL: http://genome.gsc.riken.go.jp/
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda

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S. Hashizume, W. Hayashida, K. Hirozane, T. Hori, F. Imotani, K. Inishi, Y. Itoh, M. Kigawa, I. Kawai, J. Kojima, Y. Kondo, S. Konno, H. Koya, S. Miyazaki, A. Murata, M. Nakamura, M. Nomura, K. Nomura, K. R. Ono, M. Ohtono, N. Saito, R. Sakazume, N. Sano, H. Sasakura, D. Sato, K. Shibata, K. Shiraki, R. Takagami, T. Takeda, Y. Waki, K. Watahiki, A. Muramatsu, M. and Hayashizaki, Y. Direct Submission

Comparational Analysis of Full-Length Mouse cDNAs Compared with Human Sequences. *Mamm. Genome*, 12, 678-687 (2001).

Normalized Sequence Libraries of cDNA Libraries. To prepare full-length cDNA libraries for large-scale discovery of new genes. *Genome Res*, 10 (10), 1617-1630 (2000).

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. *Genome Res*, 10 (11), 1757-1771 (2000).

FEATURES

source

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location/Qualifiers
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/strain="C57BL/6J"
/db_xref="taxon:10090"
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**BASE CO
ORIGIN**

Query Match 79.1%; Score 17.4; DB 14; Length 773;
Best Local Similarity 94.7%; Pred. No. 2.8e+03;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 CGATCGGGGCGGGCGGAGC 22
Db 147 CGAGCGGGGCGGGCGGAGC 165

Dbb 147 CGAGCGGGCGGGCGAGC 165

Search completed: February 18, 2004, 17:13:00
Job time : 1636 secs